USAAMRDL-TR-76-5

ADA035217

MM&T - PULTRUDED COMPOSITE STRUCTURAL ELEMENTS

Goldsworthy Engineering, Inc.
2917 Lomita Blvd.
Torrance, CA 90505

December 1976

Final Report

D D C
RECEIVED
FEB 4 1977
A

Approved for public release;
distribution unlimited.

Prepared for

U. S. ARMY AVIATION SYSTEMS COMMAND
P. O. Box 209
St. Louis, Mo. 63166

EUSTIS DIRECTORATE
U. S. ARMY AIR MOBILITY RESEARCH AND DEVELOPMENT LABORATORY
Fort Eustis, Va. 23604

391379

EUSTIS DIRECTORATE POSITION STATEMENT

This manufacturing methods and technology project has been accomplished as part of the U. S. Army Aviation Systems Command Production Engineering Missions Program. This report identifies the application of pultruded structural elements, provides design allowables data, and establishes a fabrication technology base. Results of this contract are being integrated with other R&D efforts at the Eustis Directorate to improve the characteristics of future Army aircraft systems.

Robert L. Rodgers of the Technology Applications Division served as technical monitor for this effort.

SECURITY CLASSIFICATION OF THIS PAGE *(When Data Entered)*

REPORT DOCUMENTATION PAGE		READ INSTRUCTIONS BEFORE COMPLETING FORM
1. REPORT NUMBER USAAMRDL-TR-76-5	2. GOVT ACCESSION NO.	3. RECIPIENT'S CATALOG NUMBER
4. TITLE *(and Subtitle)* MMT - PULTRUDED COMPOSITE STRUCTURAL ELEMENTS		5. TYPE OF REPORT & PERIOD COVERED FINAL rept.
		6. PERFORMING ORG. REPORT NUMBER
7. AUTHOR(s) Brian H. Jones William Jakway (Appendix C - William Jakway)		8. CONTRACT OR GRANT NUMBER(s) DAAJ02-74-C-0053 new
9. PERFORMING ORGANIZATION NAME AND ADDRESS Goldsworthy Engineering, Inc. 2917 Lomita Blvd. Torrance, CA 90505.		10. PROGRAM ELEMENT, PROJECT, TASK AREA & WORK UNIT NUMBERS
11. CONTROLLING OFFICE NAME AND ADDRESS U.S. Army Aviation Systems Command St. Louis, Missouri 63166.		12. REPORT DATE December 1976
		13. NUMBER OF PAGES 123
14. MONITORING AGENCY NAME & ADDRESS*(if different from Controlling Office)* Eustis Directorate U.S. Army Air Mobility R&D Laboratory Fort Eustis, Virginia 23604		15. SECURITY CLASS. *(of this report)* Unclassified
		15a. DECLASSIFICATION/DOWNGRADING SCHEDULE

16. DISTRIBUTION STATEMENT *(of this Report)*

Approved for public release; distribution unlimited.

17. DISTRIBUTION STATEMENT *(of the abstract entered in Block 20, if different from Report)*

18. SUPPLEMENTARY NOTES

19. KEY WORDS *(Continue on reverse side if necessary and identify by block number)*

Pultrusion	Post Forming Graphite/Epoxy Pultrusions
Composites	Pultrusion Design Allowables
Graphite/Epoxy	

20. ABSTRACT *(Continue on reverse side if necessary and identify by block number)*
The objectives of this program were to establish the manufacturing technology base in order to pultrude with a range of resin matrices and reinforcing forms, to investigate the post-forming capability of epoxy matrix pultrusions, and to derive a representative range of static and fatigue mechanical property design allowables. Additionally, structural efficiency studies were performed to evaluate the potential advantages of using pultruded substructural elements.

DD FORM 1 JAN 73 1473 EDITION OF 1 NOV 65 IS OBSOLETE

SECURITY CLASSIFICATION OF THIS PAGE *(When Data Entered)*

20. ABSTRACT - Continued

It was concluded that even using existing epoxy systems,
pultrusion is a highly viable process that results in struc-
tures possessing good mechanical properties at potentially
low cost.   Post-forming operations may be readily carried
out, which further enhances the potential application areas
for the technique.

PREFACE

TABLE OF CONTENTS

6

LIST OF ILLUSTRATIONS

7

LIST OF ILLUSTRATIONS (continued)

8

LIST OF ILLUSTRATIONS (continued)

9

LIST OF ILLUSTRATIONS (continued)

10

LIST OF TABLES

11

INTRODUCTION

Over the past seven years, a large number of research and development programs have been conducted by the Government and private industry to study the potential of advanced composites for use in aircraft structures. While these studies have shown that the use of high-stiffness, high-strength composites (such as graphite-epoxy) can reduce structural weight, widespread application has not generally occurred. The high cost of the basic fibers has been only partly responsible for this; another reason has been the generally uneconomic costs of structural fabrication. It is being increasingly recognized that the use of advanced composites will only become viable provided consideration is given to developing cost-effective manufacturing procedures which minimize the labor content inherent in producing the structure.

One production technology that holds considerable promise for reducing fabrication costs to a realistic level is that known as pultrusion - the forming of structural shapes by pulling preimpregnated oriented filaments through a heated die (Figure 1). Pultrusion is one of the few basic processes in the reinforced plastics industry that converts primary raw materials directly into a finished product, continuously and automatically.
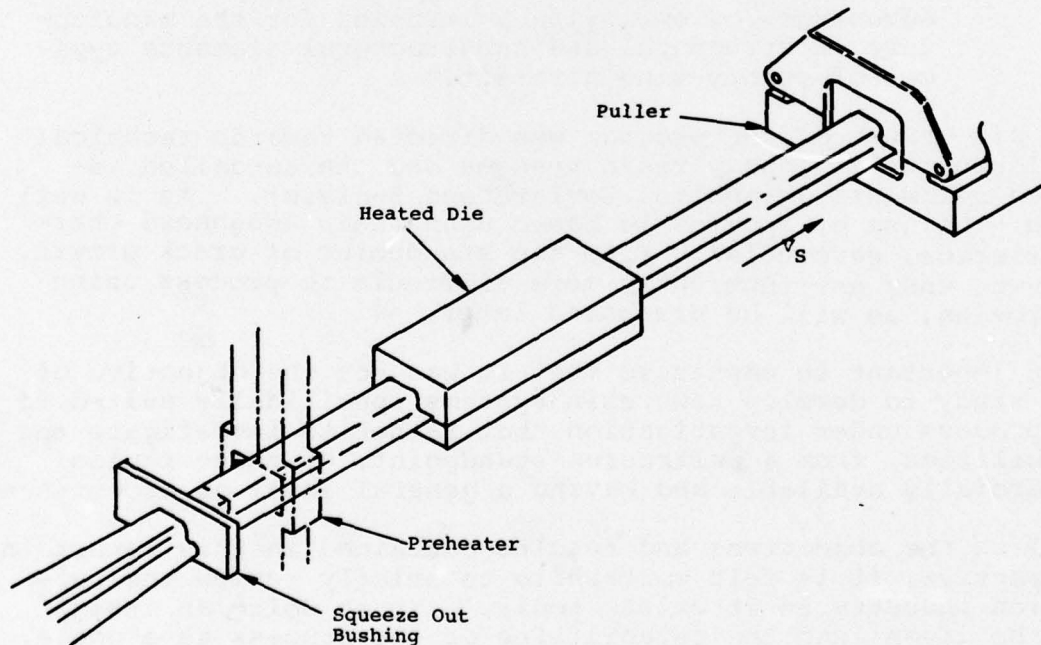
Figure 1. Pultrusion Process.

13

For some time, it has been recognized that the use of pultruded composite elements in aerospace structures offers the potential for realizing highly cost-effective designs.[1] This is particularly evident in the case of rotary-wing vehicles, where the generally favorable geometric and loading conditions suggest many design concepts highly suited to a pultruded approach. However, in order that adequate designs can be established, a data base of material properties must be available and the processing conditions must be capable of realistic definition. The purpose of this program was to provide such information.

Investigations were conducted in three phases:

1. To define the fabrication technology base for the pultrusion of cured and partially-cured structural shapes with the objective of enhancing the design potential of the pultrusion technique for a variety of structural configurations.

2. To provide initial design allowables data (static and dynamic) for pultruded structural materials to form a base for a design philosophy and criterion.

3. To identify the potential economic and performance advantages of employing pultrusion for the manufacture of structural and substructural elements typical of rotary-wing aircraft.

The main thrust of the program was directed towards technical developments with epoxy resin systems and the so-called advanced filaments (graphite, Kevlar® and S-glass). As is well known, the use of epoxies is based upon their toughness characteristics, particularly from the standpoint of crack growth. However, they are inherently more difficult to process using pultrusion, as will be discussed later.

It is important to emphasize that it was not the objective of this study to develop new resin systems specifically suited to the process under investigation, but rather to investigate the capabilities, from a pultrusion standpoint, of those resins commercially available and having a general level of acceptance.

To place the objectives and results contained in this report in perspective, it is felt worthwhile to briefly review the pultrusion industry as it exists today. In so doing, an insight will be given into the capabilities of the process as a whole.

GENERAL DESCRIPTION OF THE PULTRUSION PROCESS

The pultrusion process can be performed by up to 12 machine type variations,[2] characterized by

(a) intermittent or continuous pulling

(b) horizontal or vertical mounting

(c) tractor or reciprocating - clamp pullers

(d) conductive, dielectric or inductive curing (or combinations thereof.)

The thermosetting matrix is polymerized by the addition of heat, which may be added prior to entering the die in addition to during passage through the die. When heat is added during both periods, the cure is described as being augmented.[3] The die material is generally made of suitably plated steel, but fluoroplastics and ceramics may also be used.[3] With other than dielectric or inductive preheating, production rates are ultimately set by the thermal conductivity of the resin system.[4] Suitable preheating systems can increase production rates significantly, the limits being set by the onset of thermal cracks due to mass effect and degraded surface finish.

It is often erroneously assumed that pultrusion employs only unidirectional reinforcements. This is not the case; almost every accepted form (roving, tow, mat, cloth, braiding), from all types of filaments (glass, graphite, Kevlar, boron, steel, nylon, etc) may be used. Consideration must, however, be given to:

(a) intrinsic strength and stiffness of the impregnated material (adequate axial strength must be present to sustain instantaneous pulling forces.)

(b) thermal conductivity and specific heat (second order-effects, influencing rate of heat transmitted from die, etc.)

(c) volume ratio of reinforcing-to-matrix (controls mass effect during curing.)

If some form of dielectric or inductive preheating (injection of heat prior to final cure in the die) is being used, the electrical properties of the filaments (dielectric constant, loss tangent, etc.) become significant.

A pultrusion machine having continuous pulling capability

15

employing reciprocating clamp pullers together with augmented cure using radio-frequency dielectric augmented cure is shown in Figure 2.

The material properties required in the finished product can dictate that angle plies of fibers are necessary. Fibers oriented in the direction of pulling are best against the die surface. A cloth pulled in the warp direction will have the fill yarns displaced due to the friction of the die. The displacement will not normally be sufficient to cause jamming in the die, but continuous filament mat with a binder of low solubility in the resin can be used against the die surface. When other than random (mat) or 0°, 90° (cloth) fiber orientations are required, then braided fabric tapes can be used. Since they are fabrics, the orientations are balanced, i.e., ± 20°, ± 45°, etc.

The general requirements of the pultrusion process can be combined in a variety of ways to provide an economical solution to the fabrication of composite products while maintaining consistent quality, material properties and high production rates.

PROCESS CONSIDERATIONS

The process of pultrusion is concerned essentially with:

(a) ensuring that the resin matrix is subjected to a temperature-time history sufficient to ensure an adequate level of polymerization prior to being gripped by the puller system. At the same time, the combined effects of thermal and mechanical stress gradients caused by heating, cooling, shrinkage and pulling must not cause unacceptable material property degradation.

(b) ensuring that the impregnated material, during passage through the die, does not create shear forces that give rise to pulling requirements in excess of the intrinsic strength of the part or that cause surface spalling.

Obviously, (a) is a function of the level of preheat (if augmented cure is being employed), resin reactivity (due to hardeners, accelerators, and catalysts), die temperature, die length, stock speed, stock dimensions, mass effects and stock transverse thermal conductivity. Shear forces arise from:

1. Viscous shear - a function of temperature and the amount of diluent used. As the impregnated mass enters the heated die, the viscosity first drops and

16
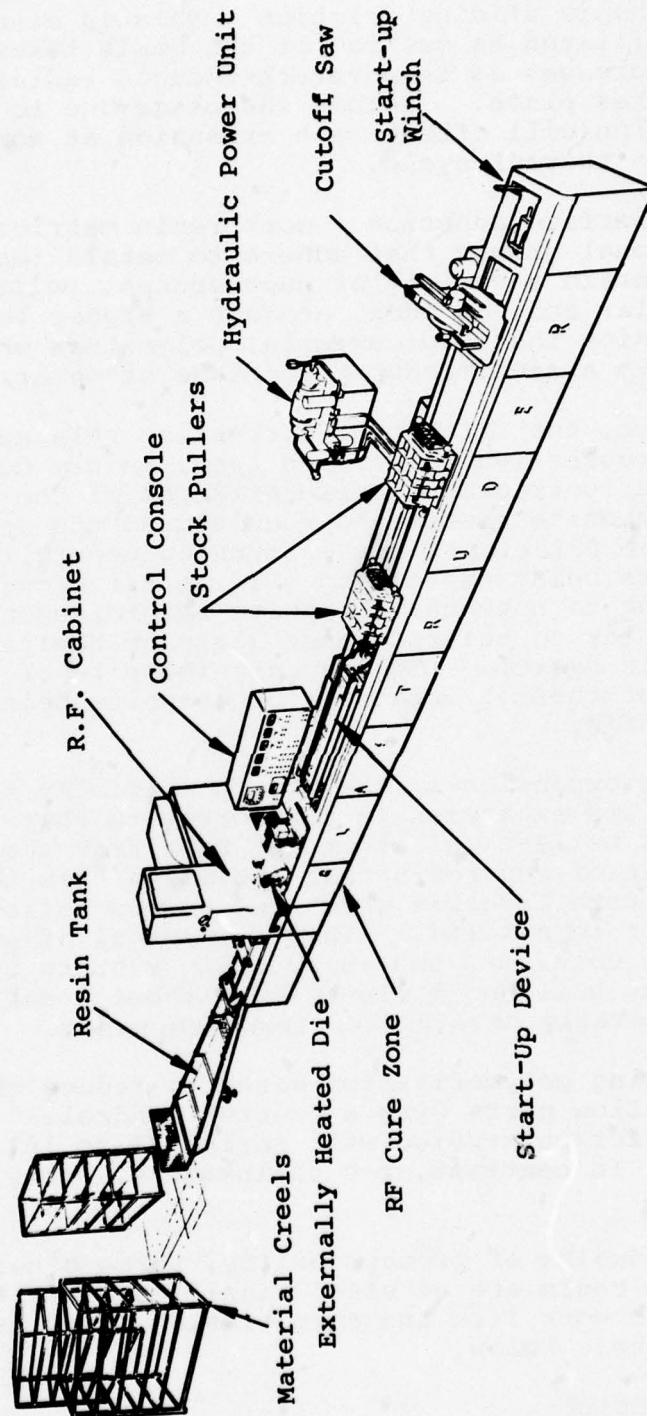
Figure 2. Typical Pultruder Using Augmented Cure on Multiple Product Streams.

Resin Tank

R.F. Cabinet

Control Console

Stock Pullers

Hydraulic Power Unit

Cutoff Saw

Start-up Winch

Material Creels

Externally Heated Die

RF Cure Zone

Start-Up Device

E X T R U D E R

17

then rises again as polymerization proceeds.

2. Dynamic sliding friction - this is assumed to be initiated as gelling of the resin takes place and increases as temperature-induced radial expansion takes place. Volume shrinkage due to polymerization will offset such expansion at some point in the thermal cycle.

3. Interface adhesion - most resin matrices have functional groups that adhere to metals (epoxy molecules contain a variety of such groups, polar and non-polar and, as such, produce a higher level of adhesion than, for example, polyesters which do not have a significant percentage of polar groups).

As noted above, the frictional forces are related to the level of radial stresses (which are, in turn, mainly dependent upon the expansion/contraction characteristics of the resin), the effective composite restrained modulus, and the instantaneous coefficient of friction. Since pultrusion generally results in some filaments being exposed to sliding frictional effects, it is interesting to note that graphite reinforcements are fundamentally easier to pultrude than glass or Kevlar when using the same resin system. This occurs in spite of the transverse coefficient of thermal expansion of graphite being of the order 20×10^{-6} in/in/°F.

Resin thermal expansion in the die is initially relieved by backflow. The expansion in the composite that increases die forces occurs between gellation and exit from the die. The characteristic of the resin that minimizes this effect is a sharp temperature level at which gel occurs followed by rapid polymerization to a solid. This is typical of polyester resin systems. In contrast, the epoxy resin systems typically exhibit a steady buildup in viscosity without a definite gel point and generally have slower reaction rates.

Shrinkage during polymerization works to reduce the die forces, except for hollow parts with a central mandrel. Typical polyester resins for pultrusion will shrink 7% to 10% during polymerization. In contrast, the shrinkage of epoxy resins is 1% to 6%.

From the standpoint of processability, three other characteristics of the resin are of vital significance, viz., filament wetting, resin work life and cure time. These aspects are discussed separately below.

Viscosity Effects

During wet impregnation of the fiber stream, the viscosity of

the resin system is a primary factor in wet-out and a secondary factor in controlling resin content. Basically, wetting of the fibers is a function of the relative surface energies of the solid and liquid involved. The fiberglass and graphite fibers have finishes or surface treatments to ensure good wetting; the Kevlar® fibers generally do not. In addition, the viscosity of the resin must be sufficiently low to ensure that the wetting is not transport limited. At the same time, it must have a high enough viscosity to prevent excessive draining from the fiber-resin stream after leaving the wet-out bath.

Experience has shown that, in general, good wetting will be obtained with viscosities below 1200 centipoise and excessive drainage will not occur at viscosities above 200 centipoise. Special techniques can be employed to operate outside this viscosity range. The resin can be heated to lower the viscosity below 200 centipoise, then the fiber-resin stream cooled after leaving the bath to increase the viscosity and prevent drainage. For higher viscosities, the rate of pulling can be reduced or the bath length increased.

Above about 15,000 to 20,000 centipoise, the viscous drag of the resin at the squeeze-out bushings, guides, or the die entrance may become high enough to commence breaking the filaments at the surface of the stream.

Resin Work Life

The pultrusion process is an automated, high-production process that realizes its greatest economies through continuous operation. A resin system, to be compatible with this type of operation, should have a long pot life or gel time at bath temperatures. Then, large batches of resin can be prepared that will be usable for a protracted period of time.

A work life of 1 week is desirable, 8 hours is acceptable, and as a minimum, it can be equal to the time required to pass the resin volume in the impregnation tank through the die at operating speeds. This latter time will allow fresh resin to be added to the bath as the level is drawn down, thereby allowing continuous operation.

Cure Time

In the pultrusion process, a fast cure time permits higher production rates and/or shorter dies. There is an upper limit on pulling speed, depending upon each resin-fiber-die combination. Since the polymerization of epoxy and polyester resins is an exothermic reaction, the polymerization will continue after leaving the die. In practice, a full cure is not required before the part leaves the die. It is only necessary that the

19

resin be hard enough to maintain its shape, and possess sufficient tensile and compressive strength to resist the forces exerted by the gripping/pulling system.

RESIN SELECTION

As has been emphasized previously, it was not one of the objectives of this program to evolve the optimum resin system for pultrusion, but rather to evaluate the pultrusion capabilities of those resin systems that were commercially available and which had a reasonable level of aerospace acceptance. The latter requirement demanded attention be directed primarily toward epoxies (intended for wet impregnation and preimpregnation) and secondarily toward polyesters (liquid and solid at room temperature).

Recent experience with epoxy matrix composites has indicated a tendency to absorb water in such a manner as to subsequently affect mechanical strength at elevated temperature [5]. As a consequence, attention is being directed toward the potential of high-performance polyesters.

For each resin type, level of compatability with the filamentary reinforcements used for aerospace application, i.e., graphite, Kevlar and S-glass, were quantified. The factors evaluated included:

(a) Tendency to die adherence, assessed only from the standpoint of acceptable/unacceptable. Actual levels of adherence were not measured.

(b) Change of viscosity with temperature, time and amount of diluent.

(c) Mechanical properties, evaluated in terms of interlaminar shear strength (considered to be a sensitive indicator of intrinsic resin performance, wet-out and processing control).

POLYMERIZATION THERMOCHEMISTRY - POLYESTERS AND EPOXIES

To aid evaluation of the results to be presented, it is proposed to briefly review the thermochemistry of both polyesters and epoxies as it applies to the pultrusion process. A more detailed discussion is to be found in Appendixes A and B of this report.

Polyesters

The unsaturated polyesters generally used in reinforced plastics consist of an ester dissolved in polymerizable monomer which

provides three-dimensional chain cross-linking.

The solution is generally polymerized under the influence of heat and/or peroxide catalyst (more correctly an initiator) to produce an infusible thermoset. Under the influence of the energy imparted by the heat, the catalyst breaks down into free radicals which initiate polymerization. A chain reaction then occurs whereby molecules are caused to react and unite. The higher the temperature, the faster the rate of catalyst decomposition. The rate of polymerization can therefore be controlled by regulating the temperature and also the amount of catalyst used.

Theoretically, the reaction should proceed to the point where all unbroken double bonds of the original unsaturated mixture are reacted by the free radicals.

In fact, about 92-95% of unsaturation is converted.[6] Gelation occurs when about 35-40% of unsaturation has taken place, the gel hardening with a reduction of volume. An incomplete cure exists when a total of approximately 80% of the unsaturation has been utilized.

Epoxies

Of the three major types of epoxy in commercial use, the epichlorohydrin/bisphenol A resins are most readily suited to pultrusion. The epoxy novalacs, used frequently as the basis for prepregs, may be used for wet impregnation pultrusion processing only, with some difficulty.

In the pultrusion process, curing agents (rather than catalysts) are generally used as the hardener, amines and acid anhydrides finding the greatest range of suitability. Aliphatic amines will give tight fast cures at room temperature but usually have a short pot life and yield products with low heat deflection temperatures. Aromatic amines result in a product with good heat deflection characteristics and chemical resistance, in addition to having a reduced tendency to adhere to the die wall when compared to anhydrides. The suggested chemical reason for this is established in Appendix A.

While anhydrides yield products with high heat deflection temperatures, good chemical resistance, low exotherms, etc., their increased tendency to die adhesion is viewed as a significant disadvantage. As indicated below, the use of internal mold releases to offset such adhesive tendencies is not recommended.

USE OF MOLD RELEASES IN PULTRUSION

Mold releases are typically fatty acids or waxes that are sol-

21

uble in the liquid resin, but come out of solution during poly-
merization.   The use of these internal mold releases was eval-
uated for the pultrusion process by a two-step procedure carried
out on stainless steel, a material known to have low tendencies
for adhesion:

Step 1.   A 304 stainless steel plate was wet sanded with No.
           600 sandpaper to simulate the honed surface of a pul-
           trusion die.   Half of the plate was buffed to a
           mirror surface and then oxidized by heating to 350°F
           for 2 hours.   One-half of the honed and mirrored
           surface was coated with carnuba wax and held at 250°F
           for 2 hours to treat the surface.   A second 304
           plate was prepared by sanding and buffing to a mirror
           surface, then oxidized at 350°F for 2 hours.

Step 2.   Stainless steel tubing, 0.25 inch I.D.,was cut 24 inches
           long and one end flared.   The bore of the tubing was
           honed to remove ripples and any draw marks.   The
           tubes were mounted in a square aluminum holder for use
           as dies.

Mold Releases Investigated

Various mold releases were combined with an epoxy resin system
that had been used for pultrusion, though with problems of ad-
herence to the die.   The resin system was Ciba 6010 Epoxy with
NMA Hardener (nodic methyl anhydride) 72 phr., Ciba Acceler-
ator 062 (benzyldimethylamine) 3 phr., and Ciba Reactive Di-
luent RD-2 (1, butanediol diglycidl ether) 16 phr.   The fol-
lowing mold releases were tested by adding 2% by weight to the
resin:

    1.   Atlas Chemical Ind., Inc.
         Atmul 84

    2.   Argus Chemical Corp.
         Stearone

    3.   Glyco Chemicals, Inc.
         Acrawax C Atomized

    4.   Ram Chemicals
         Moldgard X

    5.   Axel Plastics Res. Labs., Inc.
         Mold Wiz No. INT-18E
         Mold Wiz. No. INT-18-36
         Mold Wiz. No. INT-114-NO

    6.   Contour Chemical Co.

22

LOOB 1290
LOOB 1290.2

Results Obtained Using Mold Releases

The first plate was heated to 350°F, and each of the resin
solutions containing the mold releases was dropped on the
four surface treatments.   In all cases the resin was a soft
gel in about 1-1/2 minutes and a hard gel in about 3 minutes,
showing no inhibiting action from any of the mold releases.
With the plates at 350°F, the resins were scraped to check
for release properties from the surfaces.   The release was
poorer from the carnuba wax-treated surface than from the
clean metal.   As expected, the highly polished stainless steel
surface released better than the sanded surface.   The best re-
lease was obtained with the Mold Wiz No. INT-18E.

To further evaluate the Mold Wiz No. INT-18E, the resin was
prepared with 0, 1, 2 and 3 percent mold release.   The second
plate was heated to 350°F, and spots of each of the resin
solutions were poured on the plate. After curing for 5 minutes,
the spots were scraped off and tested for adhesion and residue
left on the plates.   Almost perfect release and minimal resi-
due were left on the plate with 3% mold release.

Evaluation of the mold release under dynamic pultrusion con-
ditions was next evaluated with the 0.25-inch I.D. tube die.
The initial set of runs were made without mold release added to
the resin and using E-glass, graphite and Kevlar® fibers.

Runs were made with both Hercules A-S and Thornel 300 graphite
fibers.   There was no discernable difference between these
fibers.   They could be pultruded continuously, but after run-
ning 30 to 40 feet the surfaces showed spalling of the resin.
After a stop in the run, of only 30 seconds, it was impossible
to restart.   The addition of 3% Mold Wiz No. INT-18E to the
resin had no effect on the results and was not deemed success-
ful.

The E-glass and Kevlar® behaved the same during the trial
runs.   In both cases, without mold release, the resin adhered
to the die and surface fibers began breaking out.   After 20 to
30 feet of material had been run, the amount of fibers broken
had reduced the cross-sectional area of the rod sufficiently
so that it broke from the pulling force.   The addition of mold
release to the resin reduced the adherence of the resin suf-
ficiently to allow a continuous run.   However, the same spall-
ing and degradation of the surface was observed as with the
graphite fibers.

23

Conclusions Reached From Mold Release Tests

The results of the testing demonstrated that internal mold releases slightly improve the pultrusion characteristics of epoxy resins in concentrations of approximately 3% by weight. Use of mold releases in concentrations above 1% by weight can degrade the resin properties. Therefore, solutions to die sticking were sought by changes in the epoxy resin system and hardener used.

RESIN EVALUATION - POLYESTER

The pultrusion process has predominately used polyester resins due to their lower viscosity at room temperature, faster cures, longer pot lives, and lower cost. However, the use of polyester with graphite is typified by low mechanical properties, especially compression and shear. The interlaminar shear strength is in the range of 6,000 to 7,500 psi.

Approach

The influence of different polyester resins and a fiber size on the mechanical properties of graphite-polyester composites was evaluated by molding unidirectional bar specimens and then testing for interlaminar shear strength.

Polyester Resins Investigated

The resins selected were as follows:

1. Silmar 685 - a liquid isophtalic-styrene resin developed for pultrusion.

2. Lubrizol 24161 - a solid isophtalic-acrylic resin with about 6% shrinkage.

3. Pittsburgh Plate Glass F34-117 - a solid, crystalline, fusible, aromatic resin with about 3% shrinkage. This resin also has a 264 psi heat distortion temperature greater than 500°F.

To improve adhesion between the graphite filaments of main interest and the polyesters, the filaments were coated with a special size, as described below.

Sizing of graphite filaments for use with polyester

The size for the graphite fibers was a special clear epoxy primer BC-401, developed by Glidden to improve adhesion of unsaturated polyester laminating resins to surfaces. Hercules AS graphite fibers were used in the test. It was found that

24

pulling the fibers through a mixture of 3 parts primer to 1000 parts epoxy thinner left a very light film of the primer on the fibers. The sized fibers were left to air-dry overnight.

Testing and Results

Using filaments that were sized and unsized, two test specimens 0.80 X 0.250 X 6.0 inches were match-die molded from each resin. The resins were catalyzed with 1.5% benzoyl peroxide and cured at 300°F for 1 hour. The fiber content of the bars was 60 v/o. The specimens were cut into test coupons and tested for interlaminar shear strength, in accordance with ASTM D-2344 -72, at a span-to-depth ratio of 4. The results of the tests are presented in Table 1.

TABLE 1. INTERLAMINAR SHEAR STRENGTH OF HERCULES AS GRAPHITE FIBER WITH VARIOUS POLYESTER RESINS

| RESIN | CONTROL | | SIZED | | % of |
	ILS (PSI)	C.V. (%)	ILS (PSI)	C.V. (%)	Change
Silmar 685	7,500	5.1	9,300	6.0	+24
Lubrizol 24161	7,600	6.6	8,200	5.2	+ 8
PPG F34-117	7,800	1.6	12,000	3.2	+54

Conclusions

It will be noted that for all resins the sized fibers resulted in higher composite interlaminar shear strengths. Also, the new class of aromatic polyester resins showed a marked improvement in strength and can be expected to show major improvements in the other mechanical properties.

RESIN EVALUATION - EPOXY

Selecting a resin system involved a consideration of various chemical, physical and mechanical properties. Pultrusion requires the resin system to have low viscosity, stability at impregnation bath temperatures, fast gel time, good wet-out of the fibers, and good mechanical properties. High-performance aircraft applications require the resin system to be moisture resistant, weather well, have temperature capabilities in the 175°F to 200°F range and exhibit good toughness. Two basic groups were investigated, the first conventional liquid, the second resins formulated for prepregging.

Liquid Resins

The resins were evaluated primarily for low viscosity and good all-round properties. Also considered was the data base available on the resin regarding their properties, reactions and general use. The resins selected were:

1. Shell Epon 828 - a 11,000 to 15,000 centipoise viscosity bisphenol A - epichlorohydrin type epoxy resin.

2. Shell Epon 815 - a 500 to 700 centipoise viscosity version of Epon 828 with butyldiglycidylether diluent added to lower viscosity.

3. Shell Epon 826 - a 6,500 to 9,500 centipoise viscosity epoxy resin.

4. Ciba Araldite 6010 - a 12,000 to 16,000 centipoise viscosity bisphenol A - epichlorhydrin type epoxv resin.

5. Dow DER-332 - a 4,000 to 6,000 centipoise viscosity bisphenol A - epichlorohydrin type epoxy resin.

Other manufacturers have similar resins, but with the above there is extensive data on their performance with various hardeners, accelerators, and diluents. These resins are widely used and with the proper hardeners have good properties at elevated temperatures.

Hardeners Used With Liquid Resins

The hardeners were reviewed with respect to their ability to minimize die adherence, to reduce the viscosity of the resins, to influence gel time, to influence pot life, and to influence elevated temperature properties. The hardeners selected were:

1. Nadic methyl anhydride (NMA) - a low-viscosity liquid. The anhydride hardeners are frequently used in adhesive formulations to promote the adhesion of epoxy resins.

2. Aromatic diamine (Shell Y) - a medium-viscosity liquid used at 30 phr.

3. Aromatic diamine (Shell Z) - a medium-viscosity liquid used at 20 phr.

4. Menthane diamine (E.V. Roberts RF-45) - a low-viscosity liquid used at 22 phr.

5. Metaphenylene diamine (MPDA) - a solid that, when

26

heated to 150°F, becomes liquid and is used at 14 phr.

6. Linear polyamide (Versamid 140)- a medium-viscosity liquid used at 30 phr.

7. Polymeric diamine (Furane 9231) - a high-viscosity liquid used at 12 phr.

Compared to amine curing agents, anhydrides react at a slower rate with epoxy resins in addition to requiring higher cure temperatures and longer cure time. Due to the low solubility of the anhydrides in the resin, special techniques are necessary to ensure uniform dispersion. While increased solubility can be obtained by increasing temperatures, this will usually result in shorter pot life. Anhydrides such as methyl nadic anhydride are liquid at room temperature and therefore exhibit improved solubility. In Europe, eutectic mixtures of anhydrides or the higher melting point dianhydrides are sometimes used for pultrusion processing.

Of the amines, the aromatics are generally preferred for pultrusion since they give a good pot life in addition to high heat distortion temperature and chemical resistance.

When using a primary amine, the reaction involves opening the epoxide ring through an electronegative attack by nitrogen to form an alcohol without formation of by-products. For the anhydride, reaction is initiated by the formation of an ester through the anhydride and an hydroxyl group present in the amine.

Any oxide present on the die surface will electronegatively attack epoxide groups and produce adhesion. Tendency to die sticking appears slightly greater with anhydrides than with amines, since the metal oxides or metal hydroxyl groups attack the epoxide ring in a similar manner to the secondary alcohol in the anhydride case.

Diluents Used With Liquid Resins

The diluents investigated were of two types: reactive and unreactive. They were selected for their ability to reduce the viscosity of the epoxy resins. Diluents tend to reduce the mechanical properties of the cured epoxy resins. Therefore, the intent was to add the smallest amount of diluent possible. The diluents selected were:

1. Butyl glycidyl ether (BGE)

2. 1, 4 - butanediol diglycidyl ether (BDE)

3. Dimethylformamide (DMF)

27

Process Evaluation - Liquid Epoxies

Evaluation of the resin involved:

(a) Investigating the level of die adherence.

(b) Determining the extent to which the viscosity of the resin was to be modified by application of heat or diluents in order to attain adequate wet out.

(c) Establishing characteristic curing rates while pultruding a circular rod.

(d) Testing the pultruded product to determine mechanical properties in terms of interlaminar shear strength, flexural strength and modulus.

The level of general process control was evaluated by determining the coefficient of variation observed in the mechanical properties data.

Die Adherence

Die adherence phenomena were investigated by control of die surface characteristics (treatment, finish) and by variation of the polymerization process involved in cross-linking the resin.

From the tests carried out into the influence of internal mold releases, it was determined that the most advantageous conditions to minimize die sticking would be attained using stainless steel, since this material does not readily form an oxide that possesses high electron attracting characteristics (high surface energy), was readily capable of being finished to produce low surface roughness under protracted use, and would not require replating (as would be the case with a steel/chrome plated die).

The stainless steel die, 24 inches long, is shown in Figure 3, and was designed with the objective of enabling specimens of various widths and thicknesses to be pultruded. The die, when pinned and bolted together, can form cavities with thicknesses of either 0.08, 0.10 or 0.125 inch and widths of 0.25 or 0.50 inch. The cover plates were 304 stainless steel and the shim spacers were steel chrome plated. A hole drilled into the upper cover plate to the centerline of the die accommodated a thermocouple probe that was attached to a temperature controller. After polishing, the die surfaces were conditioned by holding them at 350°F for 4 hours. Following this, one end of the die was plugged then filled with molten carnuba wax and held at 250°F for 24 hours.

28
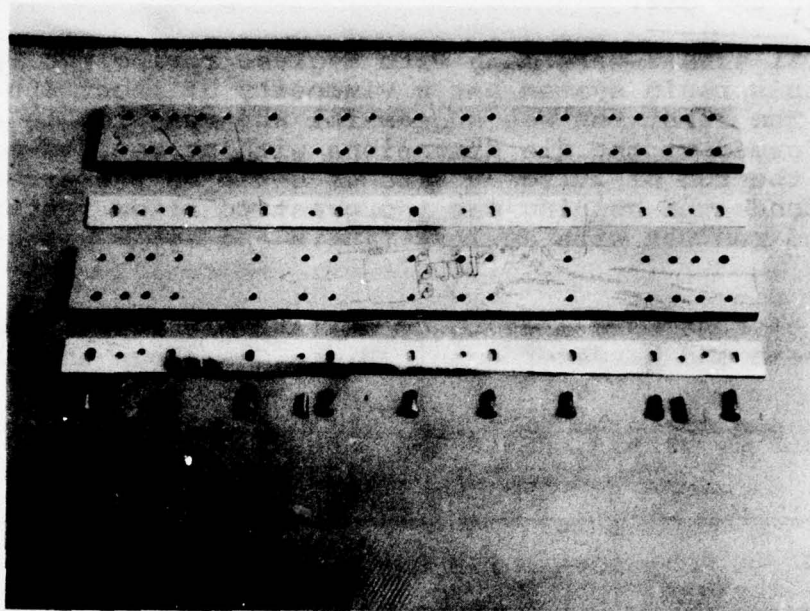
Figure 3. Break-Apart Adjustable Stainless
Steel Die - 24 Inches Long.

Four 1000-watt strip heaters were attached to the cover plates,
two on top and two on the bottom.

Initial trials through the die with E-glass roving and the Ciba
6010/NMA resin system all resulted in die adherence and termin-
ation of the run. The die was opened after the runs and it
could be seen that a resin buildup was occurring on the die
surface, about two-thirds of the way from the entrance. By
opening the die immediately following a run, while the die was
still hot, it could be seen that a resin buildup was occurring
at the point of initial gellation of the resin. The resin
buildup reduced the die opening and caused the product to jam
in the die.

The second approach was to vary the type of hardener. As re-
viewed earlier, the anhydride hardeners react by activating the

29

hydroxyl groups on the epoxy resin and proceed in a chain re-
action.   The activated hydroxyl groups, at the mold surface,
can promote bonding.   Conversely, the amine hardeners react
with the terminal epoxide groups.   The result is less tendency
to bond to the die.

The initial trials were made with E-glass roving and Epon 815/
RF45 as this resin system has a viscosity of about 600 centi-
poise.   The first run was successful and resulted in a part
that conformed to the die dimensions without rounded corners.
Stops in the run of 20 to 30 seconds could be restarted, but
at 60 seconds the bonding was too great to allow restarting.
The second run was with Kevlar® fibers and again was success-
ful.

A series of runs were made to quantitatively evaluate different
fiber, resin and hardener combinations, as shown in Table 2.

TABLE 2.   FIBER, RESIN AND HARDENER COMBINATIONS
EVALUATED FOR DIE ADHESION CHARACTERISTICS

RUN NO.	FIBER	# OF ROV- INGS	FIBER V/O	RESIN	HARDENER	DIE TEMP. °F
1	E-Glass	5	60	Epon 815	RF-45	400
2	Kevlar®	5	60	Epon 815	RF-45	400
3	Herc.-AS	15	55	Epon 815	RF-45	400
4	Herc.-AS	15	55	Epon 815	Shell Z	350
5	E-Glass	5	60	Epon 815	RF-45	400
6	E-Glass	5	60	Epon 828	RF-45	400
7	E-Glass	5	60	Epon 815	Shell Z	350
8	E-Glass	5	60	Epon 815	MPDA	400
9	E-Glass	5	60	Epon 828	MPDA	400
10	E-Glass	5	60	Epon 828	MPDA	400
11	E-Glass	5	60	Epon 815	Furane 9231	400
12	E-Glass	5	60	Epon 828	Furane 9231	400
13	E-Glass	5	60	Epon 828	Furane 9231	400
14	E-Glass	5	60	Epon 828	Shell Z	350

Run No. 5 was made with the Mold Wiz INT-18E added to the 815/
RF45 resin.   There was no change that could be discerned be-
tween this run and No. 1 without the mold release.   The higher
viscosity resin systems were run in a heated impregnating bath
to reduce resin viscosity to between 1000 and 2000 centipoise.

During these runs, the die temperature was set initially at 350°F, then the pull rate was adjusted until the fiber-resin stream emerging from the die was just solid. If the pull rate was then below 10 to 12 inches per minute, the die temperature was increased to 400°F and the pulling speed readjusted. If the resin system showed signs of being burned, the die temperature was reduced.

From these runs it was concluded that, in general, epoxy resin with amine hardeners could be successfully pultruded without sticking in the die with fiberglass, Kevlar® or graphite fibers.

Viscosity Investigations

The viscosity of the resin in the impregnation tank is of prime importance in controlling fiber wetting, resin content, air entrapment and voids. From the prior runs it was established that a practical upper limit on the viscosity of the resin should be about 1000 centipoise. From experience with polyester resins, the lower limit was set between 300 and 500 centipoise. A viscosity above 1000 centipoise requires additional attention from the machine operator to ensure wetting and a smooth fiber flow. Entrapped air in the fiber-resin stream is normally removed at the squeeze-out bushings and the die entrance. A viscosity below 300 or 400 centipoise allows air to reenter the fiber-resin stream.

All the epoxy resins selected for development have viscosities above 1000 centipoise, except the Shell Epon 815. However, this latter resin is identical to Epon 828 with about 15 phr butyl glycidyl ether as a diluent to adjust the viscosity to 500 to 700 centipoise. Therefore, the Epon 815 was dropped from the investigation.

The MPDA and RF-45 hardeners have the greatest effect on lowering the viscosity of the epoxy resins and give good elevated temperature mechanical properties. The viscosity of all the resins with these hardeners was above 1000 centipoise. The viscosities were then lowered by the addition of both butyl glycidyl ether (BGE) and dimethylformamide (DMF) to the resins. The results are given in Table 3. The viscosities were measured with a Brookfield Viscometer using a No. 1 spindle.

To evaluate these resin systems, a series of pultrusion runs were made with seven rovings of E-glass in a .10-x.25-inch die. The material from the run could then be cut into lengths for short beam interlaminar shear tests. The results of the interlaminar shear tests on the composite from the runs is shown in Figure 4. In reviewing the data it will be noted that with

31

TABLE 3. EFFECT OF DILUENTS ON RESIN VISCOSITY AT ROOM TEMPERATURE

| | Viscosity - Centipoise | | | | | | | |
| | Shell 828 | | Shell 826 | | Ciba 6010 | | Dow 332 | |
Diluent phr	MPDA	RF45	MPDA	RF45	MPDA	RF45	MPDA	RF45
BGE 15	600	-	-	-	510	-	-	-
BGE 10	-	-	750	-	-	400	500	-
BGE 7.5	-	-	-	400	-	-	-	-
DMF 5	-	-	-	-	800	630	-	-
DMF 2.5	-	-	650	550	-	-	-	-
DMF 2	-	-	-	-	-	-	-	525
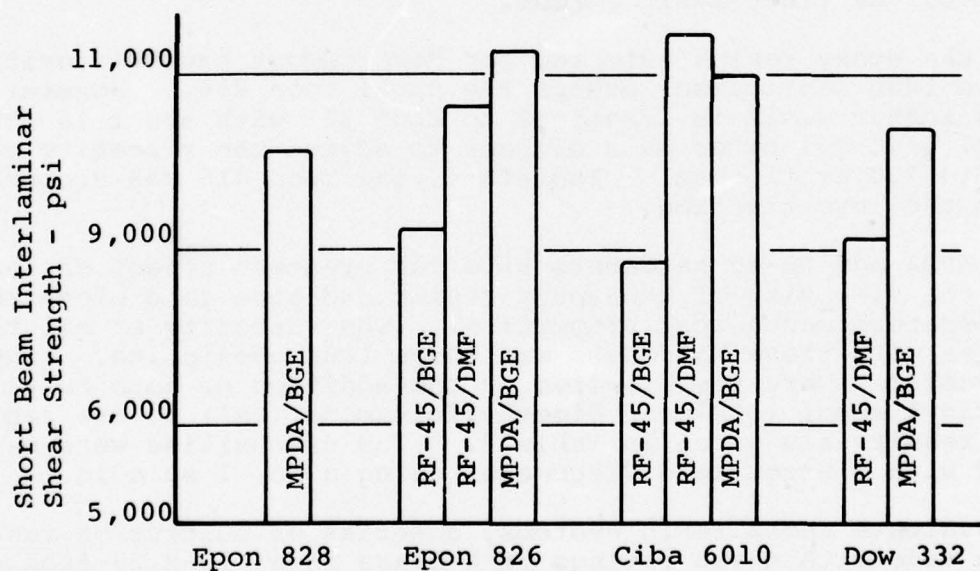
Figure 4. Effects of Various Hardeners and Diluents on the Interlaminar Shear Strength of Epoxy Resins.

the same hardener, the DMF diluent gave higher strenths, and
of the two hardeners, the MPDA gave better values.   Based on
these results, two final pultrusion runs were made using Thornel
300 graphite fibers in the 0.10-x 0.25-inch die.   To produce
60 volume percent graphite, 88 tows of the 3000 filament tows
were required.   Two resin systems were chosen for these runs as
follows:

1.	Shell Epon 826	100 parts
	MPDA	14 phr
	DMF	5 phr
	Viscosity @ 100°F	500 cps
2.	Ciba 6010	100 parts
	MPDA	14 phr
	DMF	7.5 phr
	Viscosity @ 100°F	500 cps

The runs were made with a heated and thermostatically controlled
impregnating tank maintained between 100°F and 120°F.

The results of short beam interlaminar shear tests on the graph-
ite-epoxy material from the runs were almost identical, with the
Epon 826 system being 11,700 psi and the Ciba 6010 system being
11,400 psi.   The respective coefficients of variation were
7.6% and 3.3%.   These results, in themselves, were not considered
sufficiently conclusive to enable resin selection to be confident-
ly made.   Since, in general, the addition of diluents decreases
the elevated temperature strength of epoxy resins it was felt
desirable, therefore, to make two additional pultrusion runs
using Hercules AS graphite fiber with 10,000 filaments per tow
for four-point flexure testing at 170°F together with inter-
laminar shear strength at room temperature.   The test results
are shown in Table 4.

TABLE 4.   FLEXURAL STRENGTH AT 170°F AND INTERLAMINAR
SHEAR STRENGTH AT ROOM TEMPERATURE (HERCULES
AS GRAPHITE/EPOXY)

Resin System	170°F Flexural strength (PSI)	Interlaminar shear strength (PSI)
826/MPDA/DMF	145,000	14,300
6010/MPDA/DMF	113,000	12,300

These tests identified the advantage of the lower diluent per-
centage possible with the Shell Epon 826 resin.   As a result
of this testing, the choice of the liquid epoxy resin system for
the program was 826/MPDA/DMF.

Tank Life

In a continuous process, the work life of the resin ideally
should be several days.   However, if the impregnating bath
volume can be kept small in relation to the resin being with-
drawn, it is practical to have work lives for the resin of the
order of 3 to 4 hours.   Less than this time causes problems in
mixing small batches and is too time-consuming.   The working
life of a resin system varies with the type of hardener, tem-
perature of the bath and mass of the sample.

To evaluate the tank life of the resin-hardener-diluent systems
studied for viscosity effects, a 1-quart sample of each of the
resin systems was held at 77°F, and the viscosity of the mixture
was tested each hour for 8 hours.   The criterion used to estab-
lish the usable   tank life was that the viscosity could not
exceed 1000 centipoise.   The viscosity was measured with a
Brookfield Viscometer using a No. 1 spindle.

The results of the test at room temperature for the Shell 828/
MPDA and Ciba 6010/MPDA, both with 15 phr of BGE, are shown in
Figure 5.
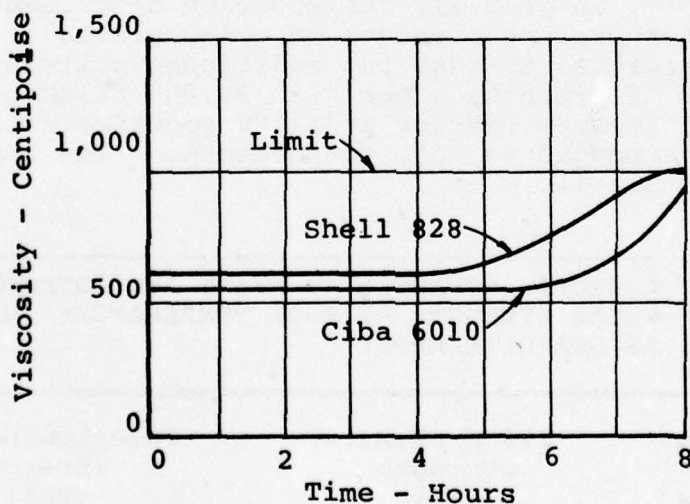
Figure 5.   Tank Life for Epoxy Resins With 14 phr MPDA
            and 15 phr BGE.

34

The results for the other resin systems at room temperature are
shown in Figures 6, 7 and 8.   The tank life of the
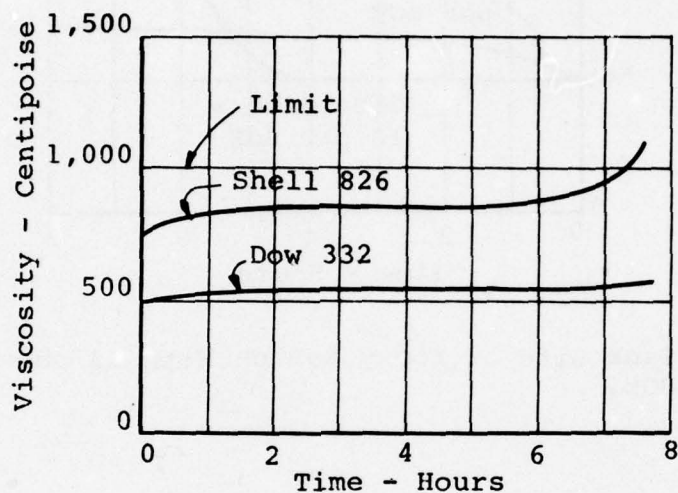
Figure 6.   Tank Life for Epoxy Resins With 16 phr MPDA
            and 10 phr BGE.

Shell 826/MPDA/DMF system, shown in Figure 7, indicates 6 hours
to reach a viscosity of 1000 centipoise at room temperature
and 4 hours at 120°F.   These items are acceptable for a pro-
duction system.
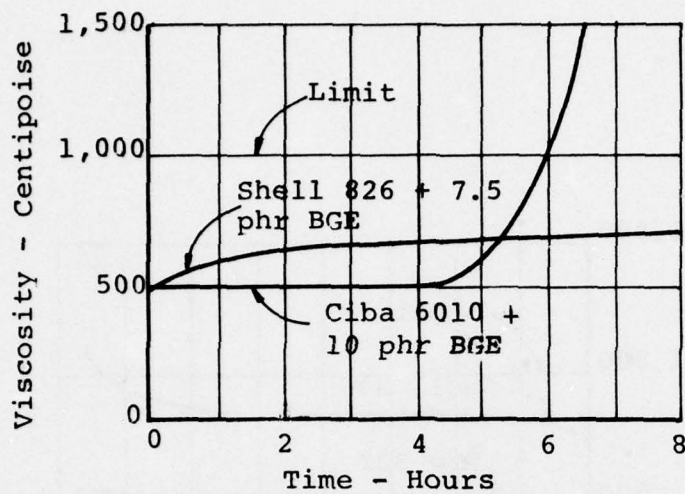
Figure 7.   Tank Life of Epoxy Resins With 23 phr RF-45 and BGE.
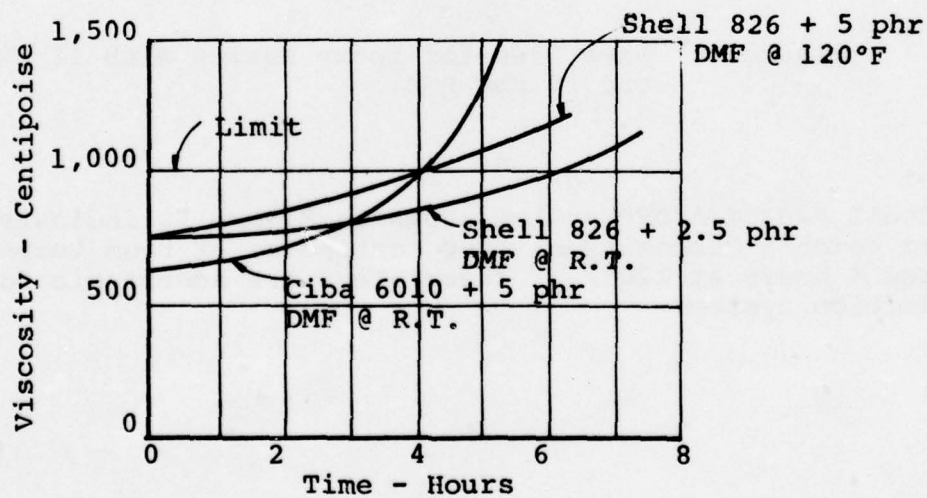
Figure 8.   Tank Life of Epoxy Resins With 14 phr MPDA and DMF

36

Cure Time Characteristics

The extremely fast epoxy resin and hardener systems that give cure times in the order of minutes also have very short work lives. As a result, this type of epoxy system is not suitable for pultrusion. The more stable epoxy systems that also are effective matrices in structural composites have extended cure times and cure schedules normally including various phases for applying vacuum and pressure to the system as well as definite temperature and time steps. (For the Shell 826/MPDA/DMF resin system, a typical cure cycle would be to hold the part at 100 psi pressure for 30 minutes at 300°F followed by removal from the mold and postcure either 1 hour at 400°F or 2 hours at 300°F.)

In pultrusion, the exposure of the resin matrix to a thermal cycle is a function of the die length and part pulling speed. Since the force required to pull a part through a pultrusion die is a direct function of die length and surface area, the practical limit to die length may be set by the tensile strength of the fiber-resin stream and the cross sectional area of the part. In this program, die length was limited to 24 inches because of the greater sticking tendency of the epoxy resins. The pulling speeds were varied from 6 inches per minute to 24 inches per minute. Below 6 inches per minute pulling speed, the part progressed in an irregular manner through the die. Therefore, speeds below 6 inches per minute were not used. The upper limit of 24 inches per minute was set by the general cure requirements of the resin.

At a die temperature of 400°F, a speed of 24 inches per minute could be obtained with the Shell 826/MPDA/DMF epoxy with any of the fibers identified earlier. The resin went from a soft gel to a hard gel in the final 8 inches of the die, giving a "cure time" of 20 seconds. By using either induction preheat with graphite-epoxy or RF preheat with fiberglass-epoxy, soft gel could be attained just inside the die entrance; under such conditions, a 24-inch die would allow a maximum pull speed of about 60 inches per minute.

The Shell 826/MPDA/DMF resin system will advance to about a 90% cure after 30 minutes at 300°F under pressure, or 2 hours at 175°F at atmospheric pressure. An additional postcure is required for maximum properties. Based on these times and temperatures, it can be appreciated that the resin coming out of the die is just advanced in polymerization past the B stage and is essentially in a hard gel state. Also, the pultrusion process can not be operated slow enough to realize a complete cure during pultrusion. Advantage can be taken of this situation where postforming of the pultruded part is desirable.

Postforming Evaluation

A series of experiments was performed on the 826/MPDA/DMF resin system to determine its potential for being postformed. Using 0.04-inch-thick by 2.25-inch-wide sheets of unidirectional graphite/epoxy, a series of tests were run to postform transverse to the filaments over radii of 2, 3.5 and 4 inches. The first attempts were made using a vacuum bag and holding the part in an oven at 175°F for one-half hour. Another set was taken through 2 hours at 175°F followed by 2 hours at 300°F, the normal cure cycle. In neither case did the part postform. By preheating the parts to 325°F to 335°F for 3 to 5 minutes, then transferring to a jig and applying pressure, the parts readily formed. It was determined that the parts had to be maintained in the jig throughout the cure cycle in order to maintain the curvature. Consistent forming could be accomplished down to a radius of curvature-to-thickness ratio (R/t) of 50 without cracking. A slight spring back was noted after release of jig pressure. Experiments performed trying to postform with radius of curvature longitudinal to the filaments were unsuccessful. Neither preheating to 325°F to 335°F prior to forming nor forming at room temperature prior to heating was successful. The specimens that were bent cold and then heated for curing experienced compressive failures on the inner surface. Specimens that were heated and bent slowly into shape could be cured successfully when R/t was greater than 250. However, on release from the jig they sprang back to an essentially flat condition.

Discussion - Liquid Epoxy Resins

From this series of experiments, an epoxy resin system has been defined that is compatible with the parameters required for the pultrusion process. The Shell 826/MPDA/DMF resin system has the following advantages:

1. Die adherence is minimum, and long runs can be made. It will permit short stoppages of 20 to 30 seconds.

2. The resin can be used with fiberglass, Kevlar[R] and graphite fibers with equal ease.

3. The viscosity of 500 to 1000 centipoise gives good wet-out of the fiber streams.

4. The tank life of about 4 hours is sufficient for continuous operation of the process.

5. The product that leaves the die is in a solid enough state to enable cutting and reasonable

handling until it is cured.

6. The 2 feet per minute pulling speed for heated
   dies is adequate. However, for parts with large
   cross sections and for higher pulling speeds, aux-
   iliary heating sources should be used to achieve
   speeds for up to 5 feet per minute.

7. Postforming transverse to the filaments is possible
   at R/t = 50 or greater for unidirectional or angle
   plied composites.

The resin system has the following disadvantages:

1. Die adherence will possibly occur for stops longer
   than 30 seconds, requiring the string-up procedure to
   be repeated.

2. While the work life is adequate under certain con-
   ditions, it is not as long as would be required for
   general usage.

3. The cure of the resin is not complete when the pro-
   duct exits the die, necessitating a postcure. This
   postcure may, of course, be carried out without fur-
   ther application of pressure or containment within
   tooling.

Process Evaluation - Prepreg Resins

As will be discussed later in this report, the suitability of
available prepreg resins was evaluated in the form of: (1) a
conventional prepreg (resin previously applied to graphite
filaments), and (2) a bulk resin (supplied in such a manner as
to inhibit polymerization by being packed in dry ice).

With the exception of Narmco Materials, no companies involved
in prepregging were willing to supply bulk resin. Through the
cooperation of Narmco, the following resins were made available
in both conventional and bulk form:

1. Rigidite 5209 - a 250°F service temperature resin
   with a fast cure cycle and 150°F liquid point.

2. Rigidite 5208 - 350°F service temperature resin
   with a slower cure cycle and 225°F liquid point.

The Narmco 5208 and 5209 resin systems are widely used in pre-
preg form for graphite-epoxy aircraft parts. As a result,

there is extensive data available on the composite mechanical
properties when cured in an autoclave.   Both 5208 and 5209
resins at room temperature have the same form, being slightly
tacky semisolids.   However, the 5209 is a 250°F service tem-
perature resin.   The 5209 melts at a low temperature and cures
faster than the 5208.

Prepreg Graphite-epoxy Tapes

Thornel 300 prepreg tapes, 3 inches wide, were obtained of both
5208 and 5209 resin.   The trials with these tapes were made
with a 0.10 x 0.25 cross section die.   The tapes were slit
into 0.50-inch-wide ribbons for feeding into the die.   It was
calculated that 8 ribbons would give a fiber volume content of
56%.   In practice, only 6 ribbons could be strung through the
die; it was planned to add the remaining 2 ribbons after the
runs were started.

The first run was with Narmco 5209.   The die was heated to
200°F and then the tape was pulled slowly till the die reached
300°F.   Simultaneously it was necessary to place a heat gun on
the prepreg entering the die to melt the resin.   In 3 trials
it was never possible to stabilize the run sufficiently to add
the additional 2 ribbons before the prepreg stream would ad-
here to the die and stop the run.   This prepreg system was
abandoned at this point.

The second set of runs was made with the 5208 prepreg.   It has
a higher cure temperature and a slower cure time.   The tapes
were again cut into ribbons, and 6 ribbons were pulled through
the die.   The 5208 resin has a gel time of about 3 hours at
300°F.   Therefore, the die temperature was brought to 350°F
before the prepreg stream was started moving slowly.   In this
case a stabilized run was achieved at 8 inches per minute and a
die temperature of 450°F.   The product stream exiting the die
was in a solid gel state.   When attempts were made to intro-
duce the final 2 ribbons of prepreg, the outer fibers commenced
shearing off at the die entrance and the fiber-resin stream
broke in the middle of the die.   After 3 attempts, the use of
prepreg was abandoned.

Actually, dry prepregs are a product form designed specifically
for another production process: lay-up and autoclave cure.
The Narmco 5208 prepreg can be pultruded by installing a temper-
ature-controlled oven system before the die to preheat the resin
to 300°F and maintain this temperature until the product stream
enters the die.   However, it would require a fairly extensive
development of the prepreg guides, bushings and controls to
achieve a production operation.

Bulk Prepreg Resins

The resins were shipped packed in dry ice and maintained at 0°F until used. The resins are tacky solids at room temperature and hard friable solids at 0°F; in the latter condition they may be easily broken into convenient sized pieces.

The initial testing was to establish the temperature at which the resins would have a viscosity of 1000 centipoise or less. Samples were heated in an oil bath until melted, and then the temperature was increased while measuring the viscosity with a Brooksfield Viscometer. The 5209 resin had a viscosity of about 900 centipoise at 200°F. However, the time required to heat the resin to this temperature was sufficient that it would gel shortly after. After several attempts to decrease the heat-up time, it was determined that there was not enough tank life, about 20 minutes, at 200°F to warrant making trial runs with this resin system.

The Rigidite 5208 resin was then melted and heated to determine its viscosity. At a temperature of 300°F the resin had a viscosity of about 900 centipoise. Checks on the tank life at this temperature showed almost no change in viscosity for 1 hour, and it required about 2 hours before gellation.

For the first pultrusion run, additional strip heaters were added to the side walls of the impregnation tank and insulation was added to facilitate maintaining a constant bath temperature. Cold resin would make wet-out difficult and excessively hot resin would shorten the tank life. The initial runs were made with Hercules AS graphite fiber and a 0.125-x 0.50-inch cross-section die using 53 tows of 10,000 filaments. The first run was made with the bath 15 inches from the die and with one squeeze-out bushing. The fiber-resin stream cooled so quickly on leaving the bath that at the squeeze-out bushing, the high viscosity of the resin began shearing off filaments. A second run with the bath 3 inches from the die was also unsuccessful. By keeping the impregnating bath exit with 0.50 to 0.75 inch of the die entrance, successful runs were made. The exit from the bath was used as the squeeze-out bushing. At a die temperature of 450°F the running speed was 15 inches per minute. The gelled parts were subsequently postcured in an oven for 1 hour at 275°F and then 2 hours at 350°F.

Interlaminar shear tests performed on the bars gave a strength of 18,500 psi and a coefficient of variation of 3.5%. This value agrees with the values reported for test specimens made from prepreg autoclave cured.

Discussion - Prepreg Resin Systems

Once the process parameters were established, the Narmco 5208 resin pultruded quite well. The short tank life of about 1 hour is satisfactory for a development program, but would preclude using the resin for a production process. In discussions with Narmco, they indicated that the resin system could be modified slightly so it would have a 900 centipoise viscosity at a lower temperature. The lower temperature would then extend the tank life. The change in the resin system would, however, result in a slight decrease in mechanical properties. Such a development may well be appropriate in further expanding the potential of epoxy resins in the pultrusion process.

FIBER SELECTION

The fiber reinforcement for pultrusion can be of any of the forms such as rovings, tows, cloth, mat, or tapes. The primary requirement of the various forms is that they be available in continuous lengths. The various forms can also be used in all combinations. Each form should have enough integrity to withstand any forces induced during pultrusion, though the more fragile types can be placed between stronger layers. In general, best results are obtained if the fibers in contact with the die surface are unidirectional rovings, tows or yarns. When cloth is against the die surface there will be a washing or displacement of the fill fibers. With some of the looser weaves the washing of the fibers can be sufficient to locally increase their density and cause jamming of the fiber stream in the die.

The unidirectional roving and tows are the most widely used reinforcements for pultruding. Yarns can also be used, but the twist makes impregnation more difficult. Also, they tend to stay as bundles of fibers with resin-rich areas between yarns. The size of the filament bundle can be very important. It requires 53 graphite tows with 10,000 filaments to fill a 0.10-x 0.50-inch cross-section die. For a part with one square inch cross section this translates into 1060 tows. While a high end or tow count can potentially offset this, too large a bundle can make adequate wet-out difficult.

As with filament winding, the filaments should have finishes applied to them to minimize the formation of "fuzz" and make handling easier. Additionally, such finishes generally promote resin wetting.

Catenaries should be absent if good fiber collimation is to be maintained and breakages from filament buildup obviated. The filamentary reinforcement should be supplied in the no-twist center-pull configuration to permit ease of splicing.

42

Woven cloths can be obtained woven or slit to given widths. These materials are convenient to enhance the transverse properties of pultrusions. In a continuous woven tape, it is only possible to get longitudinal and transverse reinforcement.

Tapes slit from mats offer in-plane isotropic reinforcement layers. With mats the fiber content is low in comparison to other forms of reinforcement.

By braiding instead of weaving, balanced angle ply tapes can be formed. The braided tapes can be obtained as either tubes or flat tape with fiber angles of $\pm20°$ to $\pm70°$. The inclusion of longitudinal yarns in the braid will prevent scissoring of the fibers during pulling.

Graphite Fibers Evaluated

During the program a number of graphite fiber tows were investigated and evaluated. In discussions with various airframe manufacturers and fiber suppliers, it became evident that the intermediate property grade of graphite was being used in the largest volume and, hence, was available at the lowest cost. Thornel 300, Hercules AS and Celanese GT30 are typical of this grade. Pultrusion runs with epoxy resin were made with all three fibers successfully. The Thornel 300 was a 3000-fiber tow and required 3-1/3 times as many packages to fill a die. The Celanese GT30 was new and not offered for sale. The Hercules AS is a 10,000-fiber tow and was supplied as four tows wound together on a bobbin. The results of short beam interlaminar shear tests carried out on samples produced using each filamentary type are shown in Table 5.

TABLE 5. INTERLAMINAR SHEAR STRENGTH OF VARIOUS INTERMEDIATE GRADE GRAPHITE FIBERS WITH SHELL 826/MPDA/DMF

Fiber	Interlaminar Shear Strengths PSI
Thornel 300	11,700
Celanese GT30	8,300
Hercules AS	14,300

43

On account of the results obtained above (Table 5), the conveniently available form of the tows, and the data base previously developed by other sources using other processes, the Hercules AS graphite fibers were chosen for this program.

Fiberglass Fibers Evaluated

Both E-glass and S-glass fibers were pultruded successfully during the program. The S-glass was Owens-Corning HTS-904 roving with 20 ends. The E-glass was Pittsburgh Plate Glass 764-NT-30 with 30 ends of K diameter fibers. This latter fiber is made specially for pultrusion. The 904 finish on the S-glass did, however, make the fiber stick in the die and as a result it was difficult to string the material prior to running. Also, the die force was higher than for the E-glass during runs. Nonetheless, since the S-glass is a higher strength and modulus fiber, it was chosen for the program.

Kevlar® Fibers Evaluated

The Kevlar® fibers manufactured by DuPont had no finish or size applied to them. As a result the adhesion of the epoxy resin to the fibers is poor. As a result the interlaminar and in-plane shear strengths, transverse tension strength, and compression strength are all lower than expected based on the fiber properties. (No temperature conditioning of the filaments was used, although this may enhance mechanical properties when using epoxy resins employing anhydride hardening agents. This fiber was included in the program because several airframe manufacturers are considering using it in hybrid Kevlar®-graphite laminates to improve resistance to low-velocity impacts.)

Braided Tape Material Used

The ability to add angle plied layers of reinforcing fibers to a pultruded product increases the flexibility of tailoring the mechanical properties of the material. For simplicity of operation and lower cost, the angle plied reinforcement should be in continuous lengths. The braiding process of weaving offers just such a capability.

To this end, Woven Structures Incorporated established the capability to braid graphite tows and fiberglass rovings into continuous flat tapes. Since the braiding process involves, of

44

necessity, sliding the tows (or yarns) over each other, a means had to be developed which prevented the graphite tow from pulling apart under the influence of these sliding forces. This was achieved by lagging the tow with a fine rayon yarn (4 turns per inch). The 904 finish on the S-glass roving produced sufficient cohesiveness to permit braiding without undue difficulty.

A +45° braided tape was made with 10,000 fiber tows of Hercules AS graphite fiber. The tape was 2.25 inches wide and had a dry thickness of 0.055 inch. To prevent scissoring of the weave during pulling, Owens-Corning HTS-904 fiberglass 20 end roving was added to the tape. These rovings ran in the 0° direction and were uniformly spaced across the tape at five rovings per inch.

A similar tape was made with S-glass roving. However, on checking the tape after manufacture, the fiber pattern was +20°. This tape was 2.25 inches wide and had a dry thickness of 0.040 inch. It also contained five S-glass rovings per inch in the 0° direction to prevent scissoring of the weave.

Both of these braided tapes were successfully pultruded with epoxy resin. Measurement of the fiber angle after pultrusion showed that in both cases the angles had only increased by 2°.

DETERMINATION OF DESIGN ALLOWABLES

The averages of the data obtained from testing the composite material produced during this program are given in the tables in this section. The material for the test specimens was fabricated by two processes:

    (a) pultrusion and oven postcure

    (b) hand lay-up, vacuum bag and oven cure

The pultruded material for test specimens included combinations of three types of fibers and two resin systems. Within each fiber-resin combination, the process conditions and fiber volume contents were held constant.

The graphite fiber was Hercules AS and it was maintained at 58 to 60 fiber volume content during all runs. The fiberglass was Owens-Corning HTS-904 roving and it was maintained at 67 to 69 fiber volume content during all runs. The third fiber was DuPont Kevlar® 49 roving and it was maintained at 67 to 69 fiber volume percent during all runs.

Process Conditions

The process conditions for the epoxy resins were maintained constant, independent of the fiber type. The Epon 826 resin system was 100 parts Epon 826, 14 phr MPDA and 5 phr DMF. The resin and MPDA were preheated to 150°F and the MPDA melted. Then the two were added together and mixed with an electric stirring motor for 5 minutes while adding the DMF. The impregnation tank was preheated to 100°F. Cooling of the resin during stirring and when added to the impregnation tank resulted in a final resin temperature of 115°F to 125°F. The bath temperature controller was reset to 130°F and the run started. The die temperature was 400°F during the runs. The fiber-resin stream speed was adjusted to just maintain solid material exiting the die. This speed was between 12 and 15 inches per minute. The pultruded material was cut on a bandsaw into 4-foot lengths and postcured in an oven for 2 hours at 175°F followed by 2 hours at 300°F.

The Narmco 5208 resin was preheated to 300°F in an oil bath which required about 1 hour. The impregnation bath was pre-heated to 300°F. Then the resin was added to the bath and the run started. The die temperature was maintained at 450°F during the runs. The fiber-resin stream speed was adjusted to just maintain a solid material exiting the die. This speed was between 8 and 12 inches per minute. The pultruded material was cut into 4-foot lengths on a bandsaw and post-cured in an oven for 1 hour at 350°F followed by 3 hours at 450°F.

Process Conditions - Vacuum Bagging

The test specimen produced by vacuum bagging involved two types of fibers and two resin systems. The fibers were Hercules AS graphite 10,000 filament tows and Owens-Corning HTS-904 fiberglass 20 end roving. The resin systems were the Shell 826/MPDA/DMF and Narmco 5208.

Two sets of specimens were made with the Epon 826 resin system; one set with graphite fiber and one set with fiber-glass. The procedure was to use the impregnating bath and squeeze-out bushing with the resin at room temperature. Then the fibers were drawn through the bath to make an impregnated tape 2 inches wide. As the tape was drawn from the bath, polyethylene fiber was placed on both sides to facilitate handling and cutting. The tape was then cut and hand laid-up on a vacuum table. The lay-ups were covered with perforated PVA film and then dry fiberglass bleeder cloth. A PVA vacuum bag was sealed to the table, a vacuum drawn and the table placed in the oven. The cure cycle was 2 hours at 200°F followed by 2 hours at 300°F.

46

For the Narmco 5208 resin, both graphite and S-glass 3-inch-wide prepreg tape were secured from Narmco. The tape was cut and laid up on the vacuum table following the same procedure as above. The cure schedule was 1 hour at 265°F followed by 2 hours at 355°F. (It should be noted that this material was cured in an oven rather than an autoclave so the properties can be expected to be lower than those generally accepted.)

RESULTS OF MECHANICAL PROPERTY EVALUATIONS

The mechanical property tests were performed using standard test specimen configurations and testing procedures as far as possible. All tests were performed on a Baldwin-Lima-Hamilton Universal Test Machine, Mark "G".

Tensile Tests

The tensile test specimens were standard IITRI coupons and are shown in Figure 9 for the 0° and off-axis fiber orientation and in Figure 10 for the 90° fiber orientation. The coupons were cut out with a diamond saw to ensure good edge cuts. The 0° and off-axis coupons were tested with Tinius Olsen rotary-type grips with horizontal closing of the wedges. The 90° coupons present a problem in that a slight offset in the grip alignment can cause a bending moment in the coupon and induce premature failure.

A special transverse tensile test fixture was built that maintained the alignment of the load application within .001 inch as shown in Figure 11.

The results of the testing are presented in Table 6. The results for the pultruded specimen tests are averages for 5 coupons.

Compression Tests

The compression specimens were 0.125 x 0.50 inch in cross section coupons and are shown in Figure 12 for all fiber orientations. Compression tests always present a problem of supporting the specimen to prevent buckling and end brooming. The load was introduced into the test fixture through a 5/8-inch-diameter steel ball mounted in two conical seats as shown in Figure 13. The coupon was held between 2 massive steel blocks, top and bottom, that gripped the coupon and introduced part of the load as shear. In this way the end load was reduced and brooming was prevented. The results of the testing are presented in Table 7. The results for the pultruded specimen tests are averages for 6 coupons and for the vacuum bag specimen tests are averages for 5 coupons.

47

were the Narmco 5208 resin, both graphite and S-glass 3-inch-
wide prepreg tape were removed from backup. The tape was man-
ufactured on the vacuum table following the same procedure
as above. The cure schedule was 1 hour at 205°F followed by
2 hours at 355°F. (It should be noted that this material was
cured in an oven rather than in autoclave so the properties
can be expected to be lower than those generally achieved.)

RESULTS OF MECHANICAL PROPERTY EVALUATIONS

The mechanical property tests were performed using Standard
test specimen configurations and testing procedures, as far as
possible. All tests were performed on a universal Test
Universal Test Machine, Model "G".

Tensile Tests

The tensile specimens were designed to conform to the
specifications for 0° and off-axis fiber orientation
and in Figure 10 for the 90° fiber orientation. The fiber-
glass polyester tabs are shown to enhance a uniform trans-
fer of the load from the grip to the specimen. The tabs
also provide a sufficient cushion of the specimen. The 0°
specimen configuration with a uniform gage length
allowed the extensometer mounting to the specimen. The
procedure followed...

A specimen mounted in the test fixture was built for
obtaining the property data. The load applied resulted in
.001 inches/sec per minute.

The test results data are presented in Table 10. The
results indicate the material properties for the average of
5 coupons...

Compression Tests

Figure 9. Tensile Test Specimens for 0° and Off-Axis Fiber Orientation.
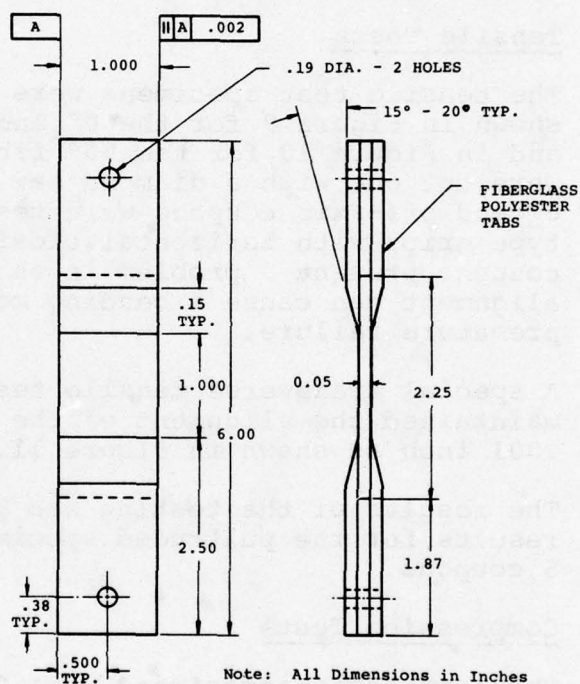
Note: All Dimensions in Inches.

Figure 10. Tensile Test Specimens for 90° Fiber Orientation.

Note: All Dimensions in Inches

48

Coupon

90° SPECIMEN

.500

2.250

.125

B   II B   .002

A   II A   .002

.500

4.380

.125

0° AND OFF-AXIS SPECIMEN

Note: All Dimensions in Inches
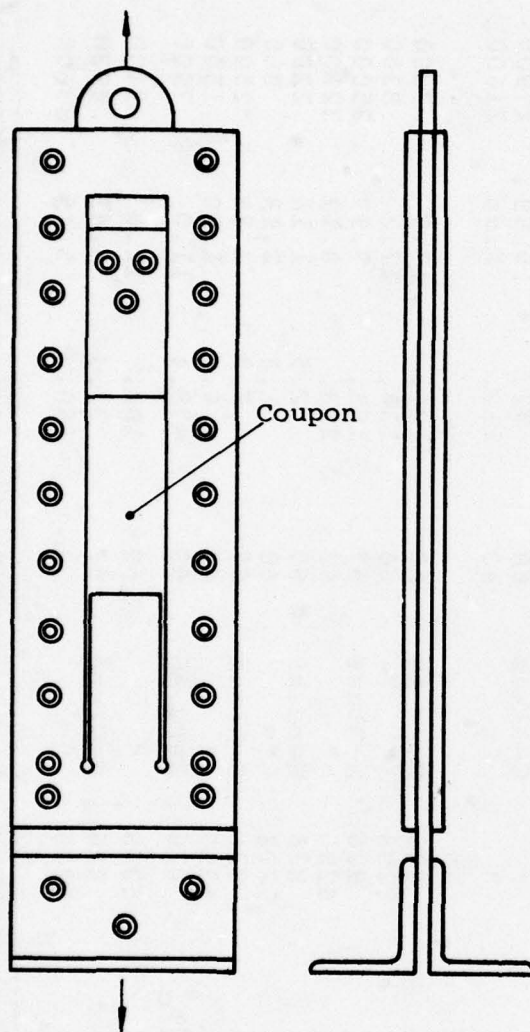
Figure 11.   Loading Fixture
to Maintain Align-
Ment for 90°
Tension Tests.
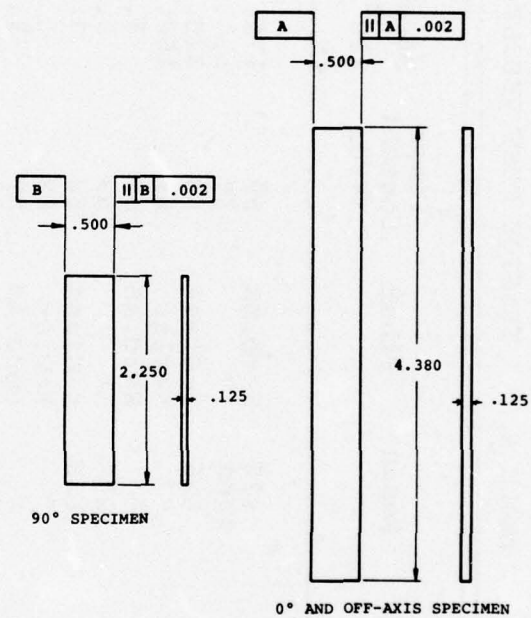
Figure 12.   Compression
Specimens.

49

TABLE 6. TENSION TEST RESULT AVERAGES

Test No.	Fiber Orientation to Load	Resin	Fiber	Fiber Content (V/O)	Strength (10³PSI)	Young's Modulus (106PSI)	Ultimate Strain (10⁻⁶in/in)	Poisson's Ratio
Pultruded								
1	(0°)c	826	Herc.AS	59	175.	20.0	8500	.20
5	"	5208	"	59	154.	19.4	7500	.18
7	"	826	S-Glass	68	228.	7.23	32800	.34
11	"	"	Kevlar	68	160.	-	-	-
2	(90°)c	"	Herc.AS	59	5.4	1.66	3700	.023
6	"	5208	"	59	2.9	2.06	1740	.029
8	"	826	S-Glass	68	7.0	1.81	2850	.094
12	"	"	Kevlar	68	1.6	-	-	-
3	(±45°)f	"	Herc.AS	59	59.	8.01	8470	.86
4	(0°±45°)s	"	"	59	93.	12.5	7400	.98
9	(±20°)f	"	S-Glass	68	92.	5.89	17900	.49
10	(0°±20°)s	"	"	68	118.	5.96	21600	.34
Vacuum Bagged								
1	(0°)c	826	Herc.AS	60	124.	17.9	7800	.32
5	"	5208	"	60	134.	17.3	8300	.25
3	"	826	S-Glass	67	152.	7.97	25000	.20
7	"	5208	"	62	213.	8.19	32300	.28
2	(90°)c	826	Herc.AS	60	2.5	1.16	2200	.097
6	"	5208	"	60	1.4	2.13	850	-
4	"	826	S-Glass	67	5.2	1.38	2500	.072
8	"	5208	"	62	4.7	1.90	1500	-
8A	(0²,±45,0²)c	826	Herc.AS	60	78.4	11.0	7900	.36
8B	"	5208	"	60	103.	12.0	9100	.48
8C	"	826	S-Glass	67	32.4	4.41	4930	.70
8D	"	5208	"	62	160.	5.45	33600	.47

TABLE 7. COMPRESSION TEST RESULT AVERAGES

Test No.	Fiber Orientation to Load	Resin	Fiber	Fiber Content (V/O)	Strength (10^3 PSI)	Young's Modulus (10^6 PSI)	Ultimate Strain (10^{-6} in/in)	Poisson's Ratio
Pultruded								
13	(0°)c	826	Herc.AS	59	96.	20.5	4700	.15
17	"	5208	"	59	133.	18.9	6800	.17
19	"	826	S-Glass	68	82.	9.3	9000	.28
23	"	"	Kevlar	68	39.	-	-	-
14	(90°)c	"	Herc.AS	59	6.4	1.4	7400	.023
18	"	5208	"	59	13.6	1.93	4400	.046
20	"	826	S-Glass	68	24.	1.64	7600	.037
24	"	"	Kevlar	68	5.1	-	-	-
16	(±45°)s	"	Herc.AS	59	25.	8.56	5700	.58
15	(0°±45°)s	"	"	59	51.	11.2	4500	.36
22	(±20°)c	"	S-Glass	68	58.	7.01	8500	.51
21	(0°±20°)s	"	"	68	66.	7.35	9400	.49
Vacuum Bagged								
9	(0°)c	826	Herc.AS	60	70.	17.2	3900	.48
13	"	5208	"	60	57.	13.1	4700	.46
11	"	826	S-Glass	67	72.	13.6	5600	.25
15	"	5208	"	62	98.	7.91	11900	.35
10	(90°)c	826	Herc.AS	60	14.	1.17	13700	.011
14	"	5208	"	60	12.	1.44	8200	.0087
12	"	826	S-Glass	67	22.	1.19	32300	.34
16	"	5208	"	62	17.	2.43	9100	.060

51

Rail Shear Test

The rail shear test specimens were 6 inches long, and when mounted in the rails they had a gage section 0.50-inch wide. To prevent rotation of the rails during loading, the rounded nose of the rails was located in a vee grove. Each coupon had 2 strain gages mounted on it in the center of the coupon. The strain gages were oriented at $\pm45°$ to the rails.

Originally, it was believed that the pultruded $+45°$ laminates would require 0° layers on both surfaces to maintain alignment of the off-axis fibers. The final braid configuration with 0° rovings internal enabled the braided tapes to be pultruded alone. This capability allowed the off-axis properties to be measured directly, rather than being derived by separating the effects of the 0° and $\pm45°$ layers by analysis.

The results of the tests are presented in Table 8. The averages for the pultruded specimens represent 6 coupons and for the vacuum bagged specimens (not shown on the table), they represent 5 coupons.

Four-Point Flexure Test Results

The flexure coupons were cut from bars 0.125 x 0.50 inch in cross section. The span-to-depth ratio was 32 and the load was applied at the quarter points, in accordance with ASTM Test Method D790. The test results are presented in Table 9.

Short Beam Interlaminar Shear Test

The interlaminar shear specimens were cut from bars with a 0.10- or 0.25-inch thickness and a 0.25-inch width. The testing procedure used was ASTM Method D-20345, with a span-to-depth ratio of 4. The test results are presented in Table 10.

Tensile Fatigue Test

The tension fatigue test was a tension-tension test conducted at constant load. The specimens were unnotched, giving a stress concentration factor of $K=1$. The stress ratio R during the test was maintained at .05. The test coupons are shown in Figure 14. The coupon is designed to be loaded primarily by a pin through the hole in end tabs and secondarily by friction from the grips. The 5.5-inch gage length was maintained the same as in the static tensile tests. During testing there was no failure of the end tabs; the majority of the failures occurred in the gage section. The cyclic rate of the testing

52

TABLE 8. IN-PLANE SHEAR TEST RESULT AVERAGES

Test No.	Fiber Orientation to Load	Resin	Fiber	Fiber Content (V/O)	Strength (10^6PSI)	Flexural Modulus (10^6PSI)
Pultruded						
25	$(0°)_c$	826	Herc.AS	59	6.7	1.49
28	"	"	S-Glass	68	3.0	.80
27	$(\pm 45°)_c$	"	Herc.AS	59	13.	2.78
26	$(0°\pm 45)_c$	"	"	59	9.5	1.98
30	$(\pm 20°)_c$	"	S-Glass	68	12.	2.29
29	$(0°\pm 20°)_s$	"	"	68	10.	2.05

TABLE 9. FOUR-POINT FLEXURE TEST RESULT AVERAGES

Test No.	Fiber Orientation to Load	Resin	Fiber	Fiber Content (V/O)	Strength (10^3PSI)	Flexural Modulus (10^6PSI)
Pultruded						
31	$(0°)_c$	826	Herc.AS	59	180.	24.2
32	"	"	Cel.GT30	60	115.	17.2
33	"	"	Cel.GY50	60	207.	16.7
34	"	"	Cel.GY70	60	93.	33.3
35	"	"	S-Glass	67	184.	7.2
36	"	"	Kevlar	68	72.	9.8
37	"	"	Thornell 300	59	181.	17.0
37A	"	5208	Hercules AS	59	210	18.4

TABLE 10. SHORT BEAM INTERLAMINAR SHEAR TEST RESULT AVERAGES

Test No.	Fiber Orientation to Load	Resin	Fiber	Fiber Content (V/O)	Strength (10^{3}PSI)
Pultruded					
38	(0°)c	826	Herc.AS	60	14.
39	"	"	Thornell 300	60	12.
40	"	"	Cel.GT-30	60	8.3
41	"	"	Cel.GY-50	60	9.5
42	"	"	Cel.GY-70	60	8.1
43	"	"	E-Glass	68	9.6
44	"	"	Kevlar	68	6.5
44A	"	5208	Hercules AS	60	11.4
44B	"	826	S-Glass	68	11.9
Vacuum Bagged					
17	(0°)c	826	Herc.AS	60	8.9
18	"	5208	"	60	6.3
19	"	826	S-Glass	67	12.
20	"	5208	"	62	6.7

.625 Dia. Ball

Test Specimen

Strain Gage

.030
Typ

Note: All Dimensions in Inches

1.50

3.50

.500

5.50

12.50

.505 DIA.

.125

Aluminum
Tab

0.125

15°-20°
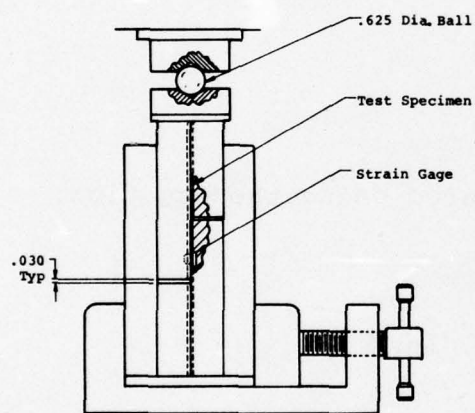
Note: All Dimensions in Inches

Figure 13. Compression Test
Fixture Showing
Load Introduction
and Supports.
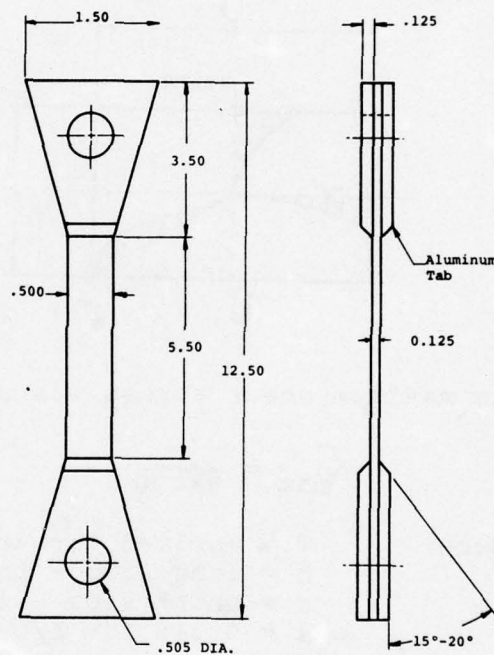
Figure 14. Tension Fatigue
Specimens.

55

was 0.5 Hertz.   The test results of the tension fatigue test
are presented in Tables 11 and 12.

Torsional Fatigue Tests

The torsional fatigue specimens were rectangular bars with a
cross section about .12 x .25 inch.   The gage length between
grips was 3.5 inches.

The shear stress distribution in a rectangular shaft is not
uniform and therefore this coupon configuration is not nor-
mally considered for torsional testing.   However, in applica-
tions such as helicopter tail rotors, a rectangular spar is a
logical choice to transmit bending and torsion loads.   Ana-
lytical solutions have been obtained for torsion of rectangular
elastic shafts. [7]   The shear stress distribution is shown
in Figure 15.
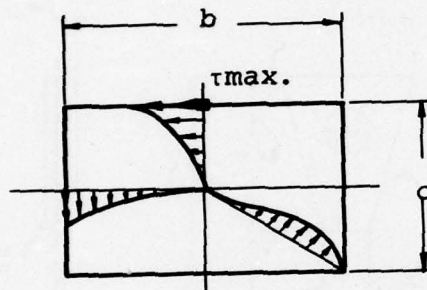
Figure 15.   Shear Stress
Distribution in a
Rectangular Shaft.

The maximum shear stress was calculated using the equation:

$$\tau_{max} = \frac{T}{8kc^2b}$$

where
T = applied torque - in. lb.
b = long side - in.
c = short side - in.
k = 0.246 for b/c = 21.

This equation is valid for a unidirectional composite since
the properties will be isotropic in the plane of the shear
stress.

The failure in a torsional test of this nature is not typi-
cally catastrophic as in the tensile fatigue test.   At the
high stress levels a visible crack actually appears; however,
at the lower stress levels it is difficult to determine failure.
For these tests, the slope of the load-deflection curve was
monitored during cycling to detect a fairly definite change in

56

TABLE 11. TENSILE FATIGUE TEST RESULTS WITH HERCULES
AS/826 RESIN, FIBER ORIENTED 0° TO LOAD –
TENSILE STRENGTH σ MAX = 175,800 PSI
K = 1   R = .05

Specimen No.	Percent of Static (%)	Test Stress (10^3PSI)	Cycles to Failure
45-1	90	156.9	1
45-2	68	119.6	1
45-3	90	156.9	4
45-4	65	114.3	3.614×10^4
45-5	80	140.6	7.774×10^4
45-6	85	149.4	1.123×10^4
45-7	"	"	15
45-8	"	"	440
45-9	"	"	310
45-10	80	140.6	2.331×10^4
45-11	"	"	490
45-12	75	131.8	130
45-13	"	"	140
45-14	70	123.1	4
45-15	"	"	12
45-16	65	114.3	1.076×10^4
45-17	"	"	No Failure in 1.889×10^6

57

TABLE 12.	TENSILE FATIGUE TEST RESULTS WITH OWENS-CORNING HTS-904 FIBERGLASS/826 RESIN, ORIENTED 0° TO LOAD – TENSILE STRENGTH σ MAX = 205,600 PSI
	$K = 1$ $R = .05$

Specimen No.	Percent of Static (%)	Test Stress (10^3 PSI)	Cycles to Failure
48-1	65	133.6	1
48-2	65	133.6	390
48-3	"	"	140
48-4	"	"	270
48-5	55	113.1	530
48-6	45	92.5	1480
48-7	"	"	700
48-8	"	"	1.25×10^4
48-9	35	72.0	1.605×10^4
48-10	"	"	1.566×10^4
48-11	"	"	7.772×10^4
48-12	25	51.4	3.9712×10^5
48-13	"	"	No Failure in 1×10^6

modulus of the composite specimens.   The point at which the
modulus changed by 10% was taken as the failure point.

The tests were conducted at a stress concentration factor
K = 1 and a stress ratio R = .10.   The test results are pre-
sented in Tables 13 and 14.

DISCUSSION OF TEST RESULTS

The Shell 826/MPDA/DMF and Narmco 5208 resin systems used in
the fabrication of the test specimens had a definite influence
on the mechanical properties of the various composite materials.
In general, the Shell 826 resin system that was formulated spe-
cifically for the pultrusion process performed best.   The
Narmco 5208 resin system was forced to conform to processing
conditions required by the pultrusion process, and as a result,
the mechanical properties reflect this condition.   The Kevlar®
fibers are still in development for use in advanced composites
and exhibited low interlaminar shear, compression and flexural
strengths indicative of a poor bond between the resin and the
fiber.

Static Properties

The longitudinal tensile properties of a unidirectional compos-
ite are dominated by the fibers.   Typical stress-strain cur-
ves for the Hercules AS graphite fiber are plotted in Figure
16.   The modulus is quite similar for either the Shell 826 or
Narmco 5208, whether pultruded (P) or vacuum bag fabricated (V).
The strengths group more distinctly by type of process, with
the pultruded specimens being significantly higher.   This
could possibly be due to inadequacies in the quality of the
vacuum specimens.

The 90° tensile properties, however, are influenced strongly
by the properties of the resin.   As shown in Figure 17, there
is a marked difference in the modulus, strength and elongation
to failure.   In this case the results from the pultruded
Shell 826 show marked superiority.

The Owens-Corning HTS-904 fiberglass behaved in a different
manner as shown in Figure 18.   The 0° tensile strength of the
pultruded 826 and vacuum bagged 5208 were essentially equal, as
were the 90° tensile tests (Figure 19).

The compression properties basically show the same trends as
indicated in Figures 20 through 23.   The results obtained
from pultrusion are higher than those obtained from vacuum
bagged specimens.   The Narmco 5208 resin tends to be better
for properties dependent on the resin properties.   The con-
trasts in interlaminar shear values are shown in Figure 24.

TABLE 13.   TORSIONAL FATIGUE OF HERCULES AS/826
UNIDIRECTIONAL COMPOSITE - STATIC
INTERLAMINAR SHEAR STRESS = 14,000 PSI,
K = 1.0, R = .10

Specimen No.	Test Stress $(10^3 PSI)$	Cycles to Failure
51-1	6.74	2150
51-2	7.31	870
51-3	6.43	150
51-4	5.35	1650
51-5	4.59	4000
51-6	4.20	4800
51-7	5.35	2800
51-8	3.82	8000
51-9	10.30	10
51-10	8.20	18
51-11	7.50	96
51-12	6.10	450

TABLE 14.   TORSIONAL FATIGUE OF S-GLASS/826
UNIDIRECTIONAL COMPOSITE - STATIC
INTERLAMINAR SHEAR STRESS =
11,900 PSI, K = 1.0, R = .10

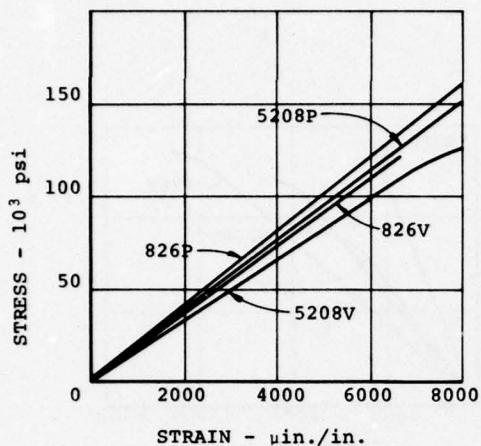| Specimen No.	Test Stress $(10^3 PSI)$	Cycles to Failure	
52-1	12.66	45	
52-2	13.14	30	
52-3	7.90	1500	
52-4	11.99	50	
52-5	8.25	1200	
52-6	7.58	1200	
52-7	7.14	2700	
52-8	6.37	5800	
52-9	6.18	6500	
52-10	11.85	96	
52-11	10.20	303	
52-12	14.80	11	
52-13	5.10	10,000	*

(* run out)

60

Figure 16. 0° Tensile Stress-Strain Relationships of Hercules AS With Various Resins.
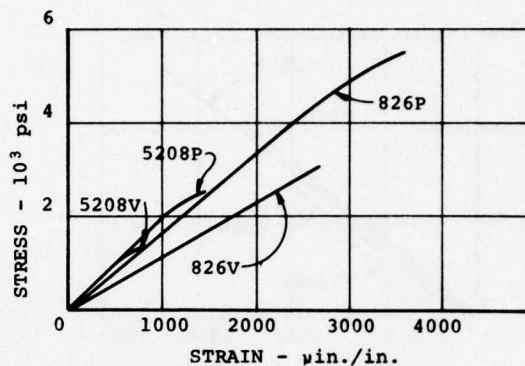

Figure 17. 90° Tensile Stress-Strain Relationships of Hercules AS With Various Resins.

Figure 18. 0° Tensile Stress-Strain Relationships of OC HTS-904 Fiberglass With Various Resins.

Figure 19. 90° Tensile Stress-Strain Relationships of OC HTS-904 Fiberglass With Various Resins.

61

Figure 20. 0° Compression Stress-Strain Relationships of Hercules AS With Various Resins.

Figure 21. 90° Compression Stress-Strain Relationships of Hercules AS With Various Resins.
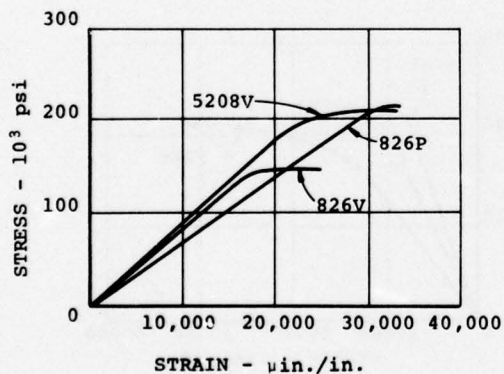

Figure 22. 0° Compression Stress-Strain Relationships of OC HTS-904 Fiberglass With Various Resins.
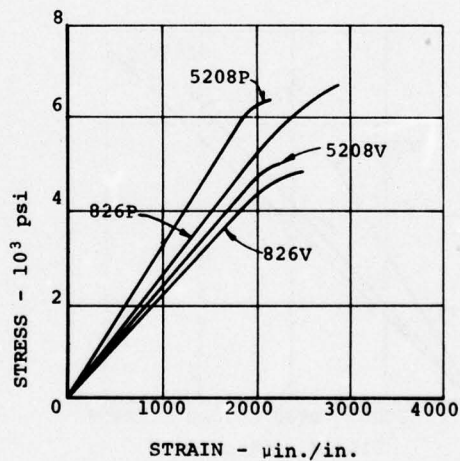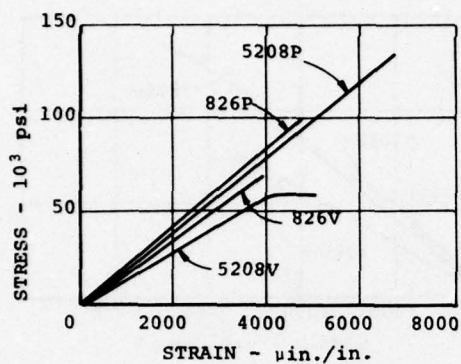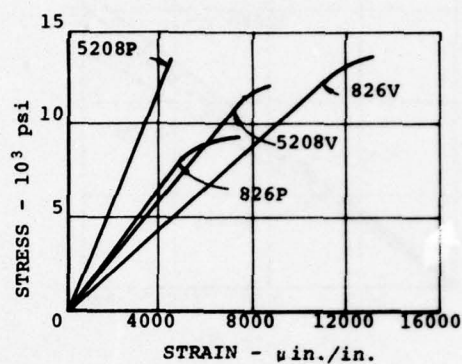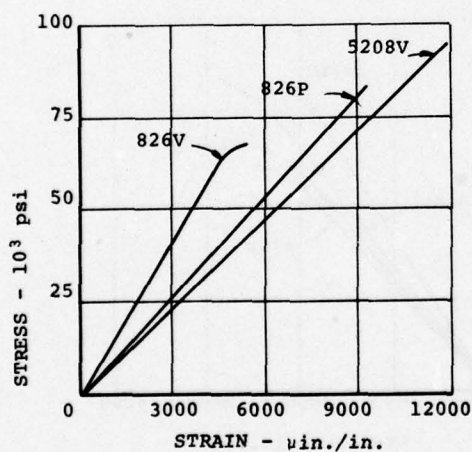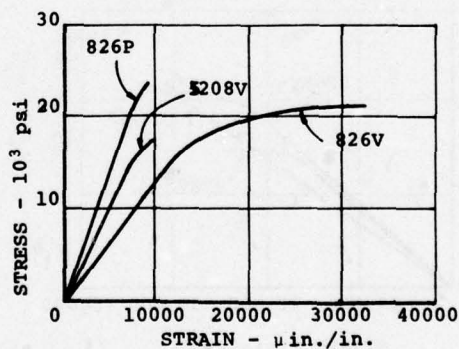

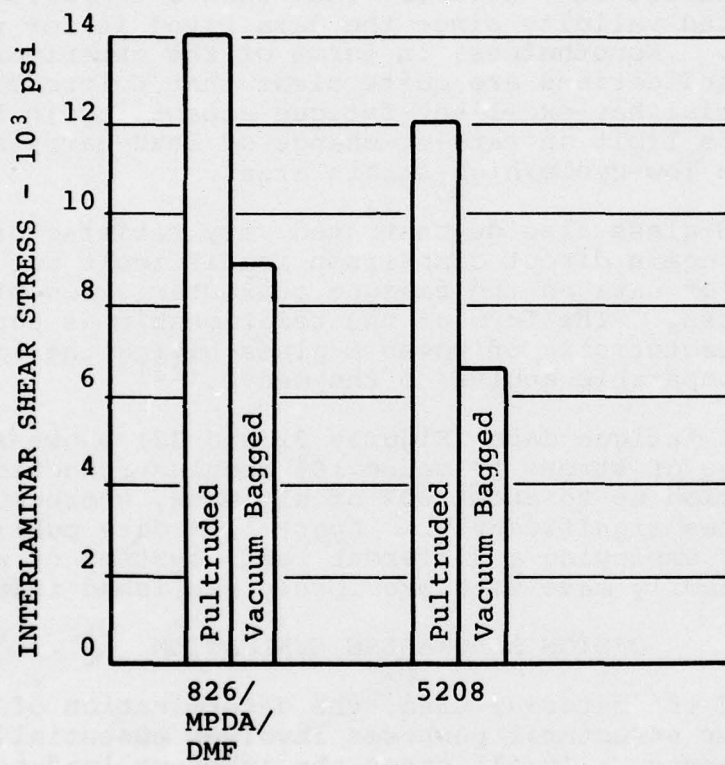Figure 23. 90° Compression Stress-Strain Relationships of OC HTS-904 Fiberglass With Various Resins.

62

Figure 24. Short Beam Interlaminar Shear
Strength Comparison.

Fatigue Properties

The S - N curves for the tensile fatigue data are shown in Figures 25 through 30. The Hercules AS/Shell 828/MPDA/DMF unidirectional laminates tested in the 0° direction had an endurance limit of 65% of the static tensile strength. This is in agreement with other published data.[8] The endurance limit for the S-glass/826 composite was 25% of the static tensile strength, which is again typical of that generally accepted.[9]

For the braided graphite materials, the data (Figures 27 and 28) compare favorably with that published in Reference 10. It must be pointed out, however, that such a comparison has only restricted validity since the data cited is for ply-on-ply material. Nonetheless, in terms of the experiments performed, the indications are quite clear that pultruded, braided material has excellent fatigue capability in terms of both endurance limit on rate-of-change of load-carrying capability in the low-cycle/high-strain area.

The braided S-glass also demonstrated very satisfactory performance but again direct comparison is difficult due to the general lack of data on the fatigue characteristics of woven, biased, S-glass. The form of the relationship is consistent with the characteristic of woven E-glass having the load applied at comparable angles to the weave.[11]

The torsional fatigue data (Figures 31 and 32) shows a high rate of change of stress to cause 10% modulus reduction with load application up to about 30% of ultimate, whereupon the rate diminishes significantly. Apart from data published in Reference 12, employing a different resin system, no comparisons can be readily made with previously published information.

DESIGN ALLOWABLES EVALUATION

Regardless of the material used, the determination of design allowables for structural purposes involves essentially the same requirements. In all cases the inherent load-carrying ability of a structure must be such that the limit load can be sustained without excessive deformation and without a significant amount of permanent set. Of equal importance is the need to ensure that the structure is capable of withstanding the operational spectrum of loading and environment for a period considerably in excess of the specified service life. These requirements do not refer to the characteristics of the material being employed; nevertheless, they are the foundations upon which the design allowables must be established. It is the unique phenomenological behavior of laminated com-

64

Figure 25.   S-N Curve For Hercules AS/Shell 826
              Unidirectional Laminate Tested in 0°
              Direction in Tensile Fatigue.
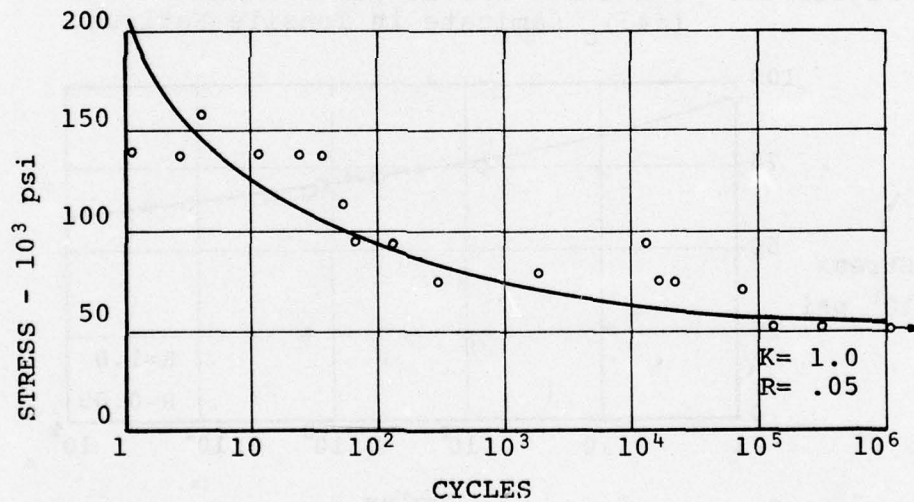
Figure 26.   S-N Curve For OC HTS-904/Shell 826
              Unidirectional Laminate Tested in 0°
              Direction in Tensile Fatigue.

65

Figure 27.   S-N Curve For Hercules AS/Shell 826
             (±45)$_c$ Laminate in Tensile Fatigue.
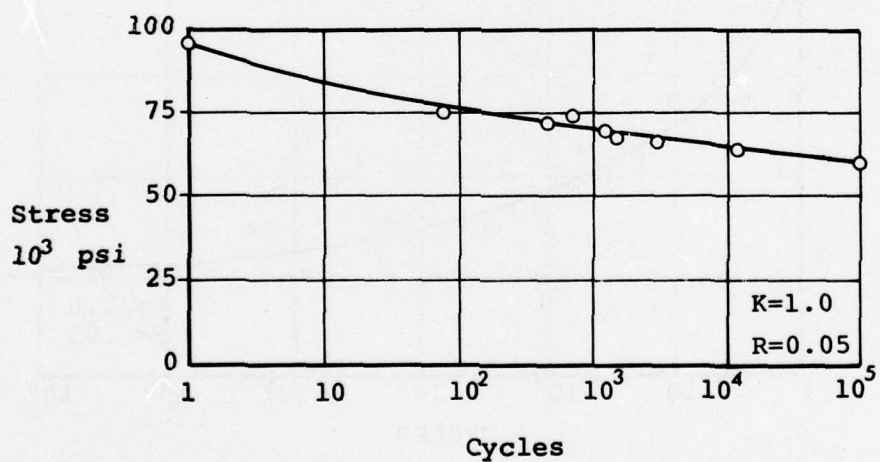
Figure 28.   S-N Curve For Hercules AS/Shell 826
             (0,±45)$_s$ Laminate in Tensile Fatigue.

66
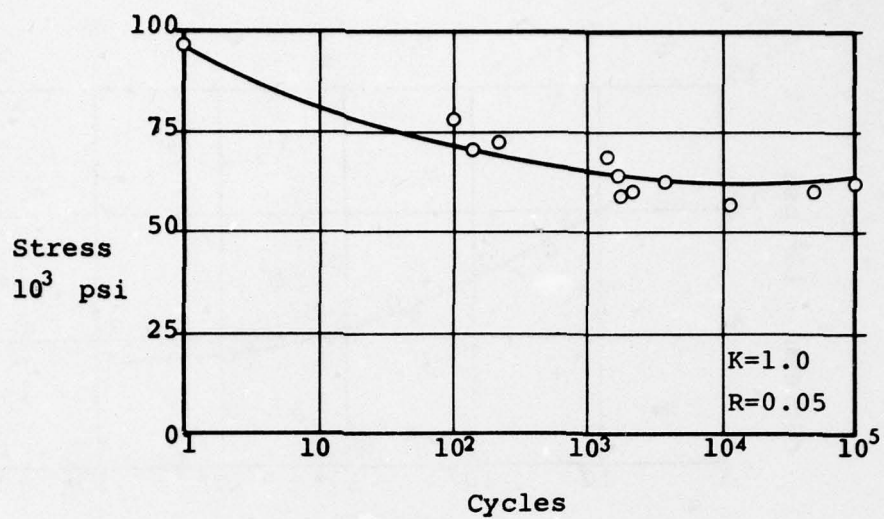
Figure 29. S-N Curve for OC HTS-904/Shell 826
$(\pm 20)_c$ Laminate in Tensile Fatigue.
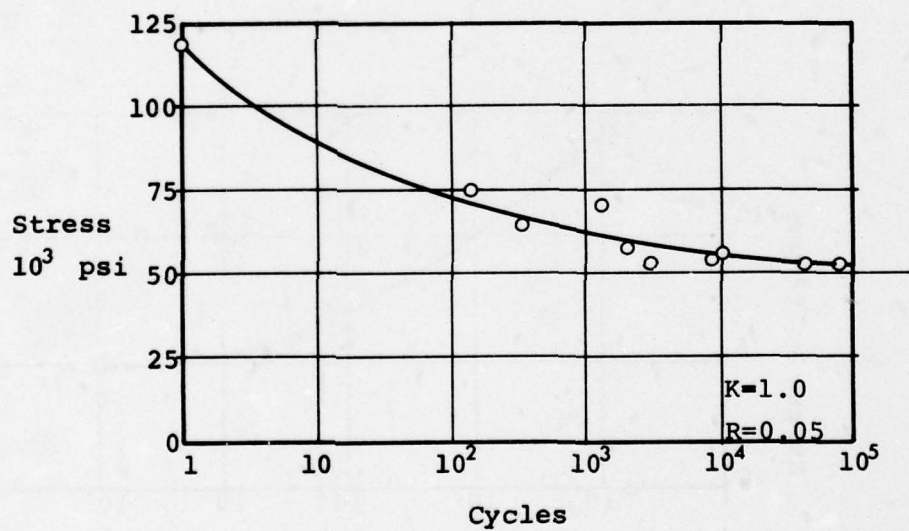
Figure 30. S-N Curve for OC HTS/Shell 826
$(0, \pm 20)_s$ Laminate in Tensile
Fatigue.
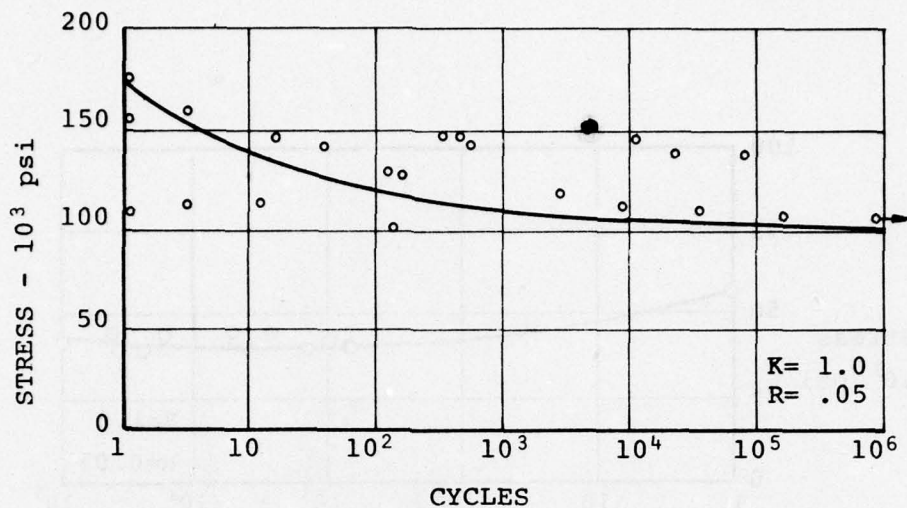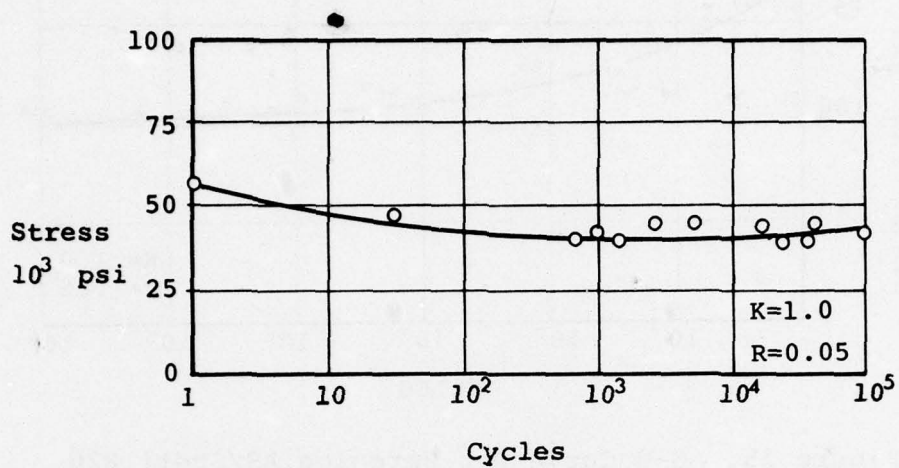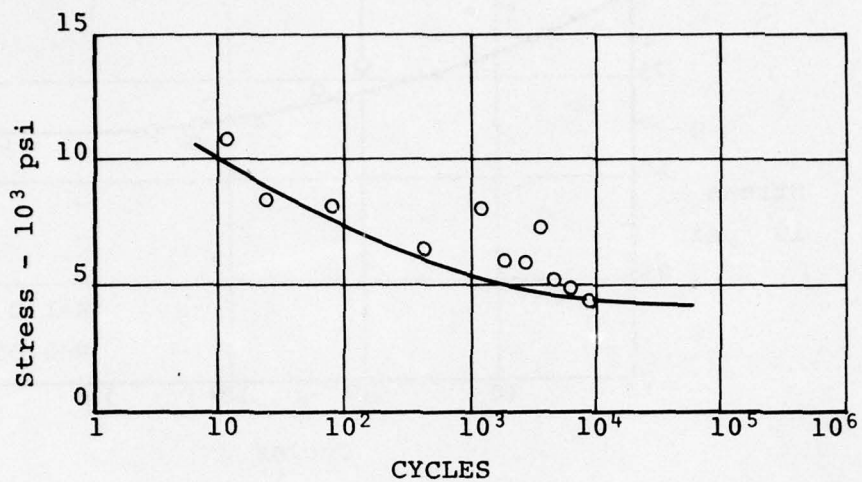
Figure 31. S-N Curve for Hercules AS/Shell 826
Unidirectional Laminate in Torsional
Fatigue.
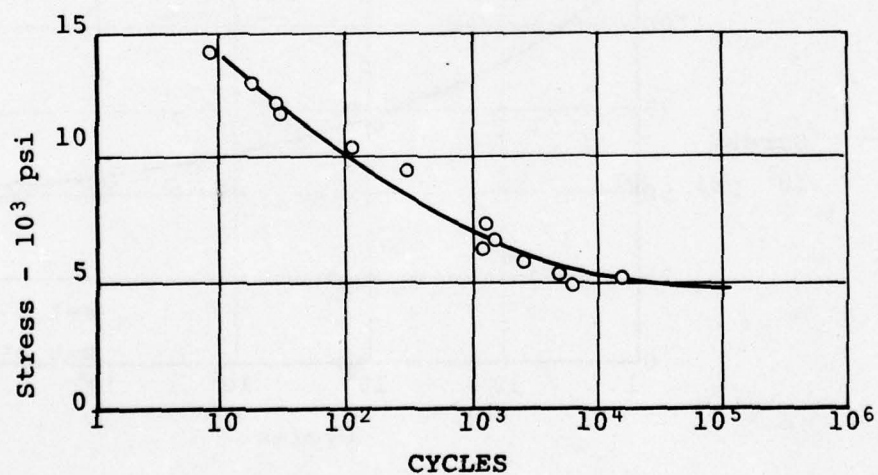
Figure 32. S-N Curve for OC HTS-904/Shell 826
Unidirectional Laminate in Torsional
Fatigue.

68

posite materials that make the determination of allowables for optimum design a challenge.

Because data are obtained from samples of finite size, it is necessary to account for variations in sample standard deviations and means for their relation to the true population by specifying confidence levels. Current aerospace practice [13] for metallic materials generally is to define a value above which at least 99 percent of the population is expected to fall with a confidence level of 95 percent (A-value). For composite materials, more realistic figures are 90 and 95 percent respectively (B-value). If the design allowable is X_{AQ} and the mean and standard deviations for the sample are $\overline{X}_s$ and σ_s respectively, then,

$$X_{AQ} = \overline{X}_s - k\sigma_s$$

where k is the one-sided tolerance factor for the normal distribution at some particular confidence level and probability as shown in Figure 33. (The significance of using data derived from small samples rather than the population is discussed in Reference 11.)

Design allowables for the Hercules AS/ Shell 826/MPDA/DMF and OC HTS-904 fiberglass/Shell 826/MPDA/DMF composite are shown in Table 15 reduced to a B basis by use of the relationship given in Figure 33. Variances are summarized in Table 15.
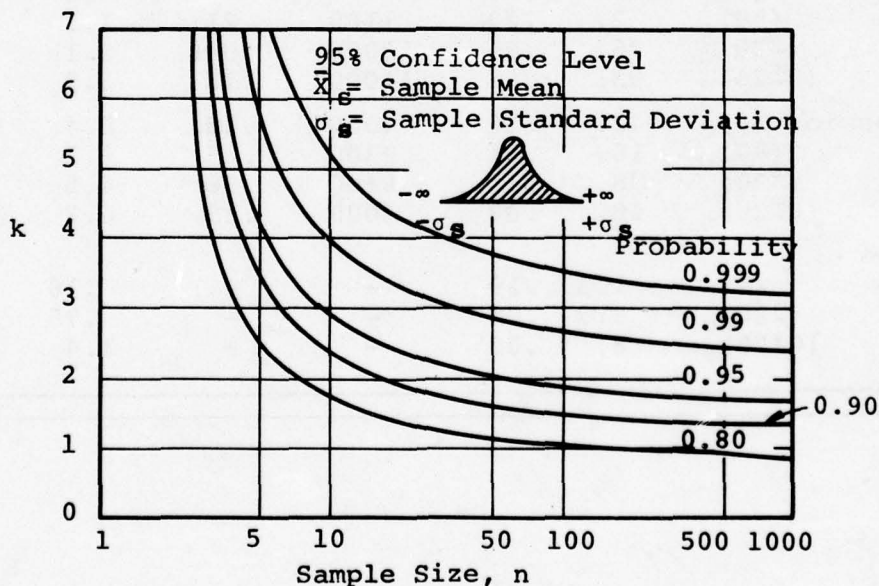
Figure 33. One-Sided Tolerance Limits and Sample
            Size for the Normal Distribution.

69

TABLE 15.	DESIGN ALLOWABLES (B-VALUE) FOR PULTRUDED COMPOSITE - SHELL 826/ MPDA/DMF RESIN

	Mechanical Property					
Load Direction to Fibers	Ultimate Strength		Ultimate Strain		Modulus	
	Ksi	C.V.	μin/in	C.V.	10^6PSI	C.V.
Hercules AS – 60 V/O						
Tension (0)$_c$	99.	.14	5500	.13	19.	.009
(90)$_c$	1.4	.32	1500	.23	.4	.29
($\pm$45)$_c$	43.	.08	2500	.28	4.4	.15
(0$\pm$45)$_s$	63.	.11	4800	.11	10.9	.04
Compression (0)$_c$	65.	.10	2600	.15	17.	.05
(90)$_c$	3.2	.16	1900	.36	.9	.11
($\pm$45)$_c$	15.	.12	1100	.35	3.4	.20
(0$\pm$45)$_s$	28.	.16	2200	.24	8.4	.09
Inplane Shear (0)$_c$	35.	.16	–	–	.6	.21
($\pm$45)$_c$	10.	.07	–	–	2.	.10
(0$\pm$45)$_s$	7.9	.05	–	–	.8	.22
OC HTS-904 Fiberglass 68 V/O						
Tension (0)$_c$	160.	.10	–	–	.5	.10
(90)$_c$	2.	.29	1100	.21	1.9	.10
($\pm$20)$_c$	76.	.05	13000	.086	5.1	.04
(0$\pm$20)$_s$	92.	.07	14000	.12	3.9	.12
Compression (0)$_c$	42.	.16	4000	.18	8.4	.04
(90)$_c$	16.	.11	2300	.25	2.1	.12
($\pm$20)$_c$	38.	.11	6400	.08	4.5	.12
(0$\pm$20)$_s$	48.	.08	4000	.19	6.2	.05
Inplane Shear (0)$_c$	1.3	.19	–	–	.16	.41
($\pm$20)$_c$	8.7	.08	–	–	.75	.24
(0$\pm$20)$_s$	8.	.06	–	–	1.4	.0

PRODUCTION AND TEST OF DEMONSTRATION ARTICLE

In order to convincingly demonstrate application of the pultrusion technology established under this program, a structural component was designed and manufactured which utilized all aspects of the results obtained. The design chosen was an integrally stiffened hybrid composite having the layup configuration and general dimensions shown in Figure 34. The panel consisted of 0° Hercules AS graphite fiber on the exterior surface and a ±45° braided inner ply of Owens-Corning HTS-904 fiberglass.

The arrangement of material used during the pultrusion operation is shown in Figure 35. To maintain precise filament placement, the ends of the impregnation tank were constructed from Teflon sheets into which a series of guidance slots were machined. These slots also served to separate the filament tows, thus aiding impregnation. The exit plate had an opening area established to control resin squeeze-out. The entrance end of the impregnation tank may be seen in Figure 36. In Figure 37 the filament resin stream entering the heated die is shown.

The pultrusion conditions were

     (a)   Resin temperature, 125°F

     (b)   Resin viscosity, 500 centipoise

     (c)   Die temperature, 350°F

     (d)   Pulling speed, 12 inches/minute

POSTFORMING OPERATION

From the 25-foot section produced, a 6-foot section was selected and postformed into a radius of 6 inches, using the forming fixture shown in Figure 38. The postforming procedure involved first mounting the parts in the fixture, followed by heating in the oven to 325°F for a period of 5 minutes. After the screws had been tightened down onto the shims, the temperature was first lowered to 200°F for 2 hours and then raised to 300°F for 2 hours for the purpose of affecting the postcure.

A section of the postformed article is shown in Figure 39.

71
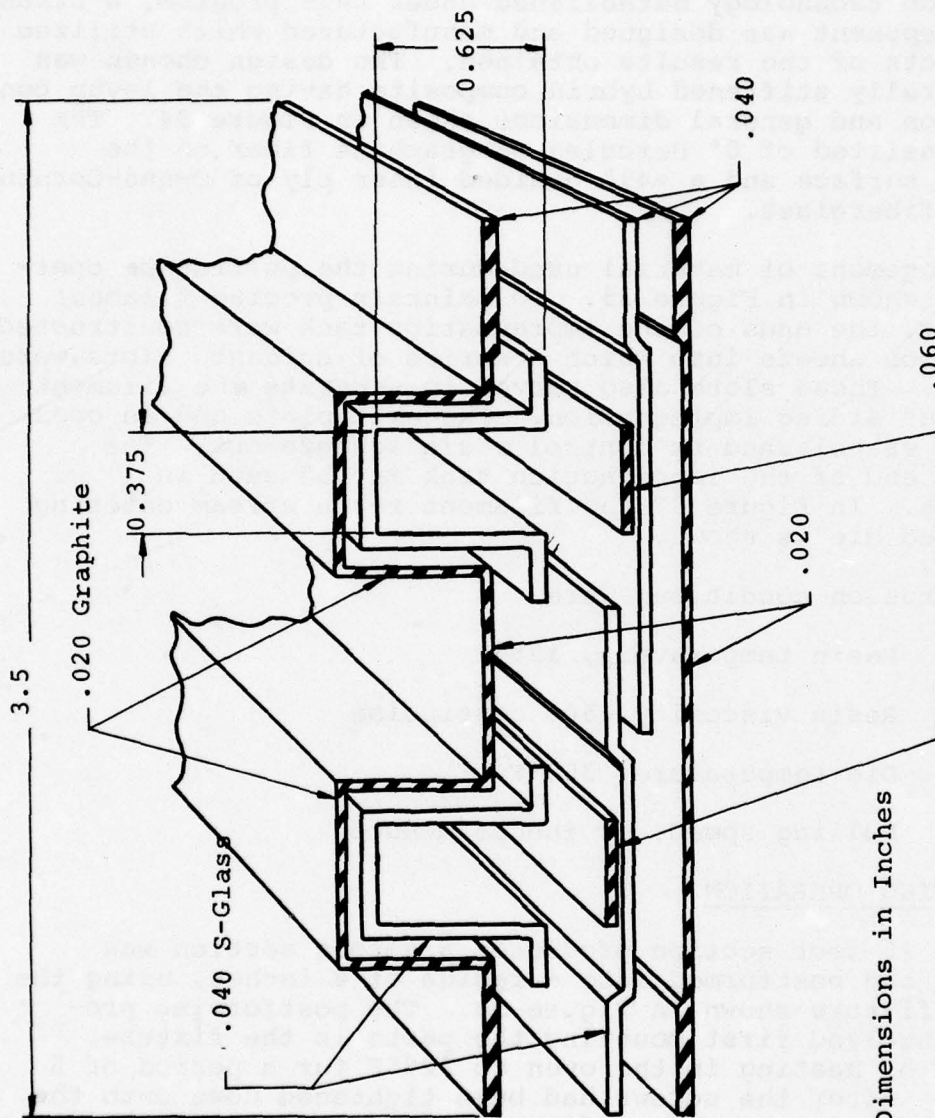
Note: All Dimensions in Inches

Figure 34. Demonstration Article - Integrally Stiffened Panel.

72
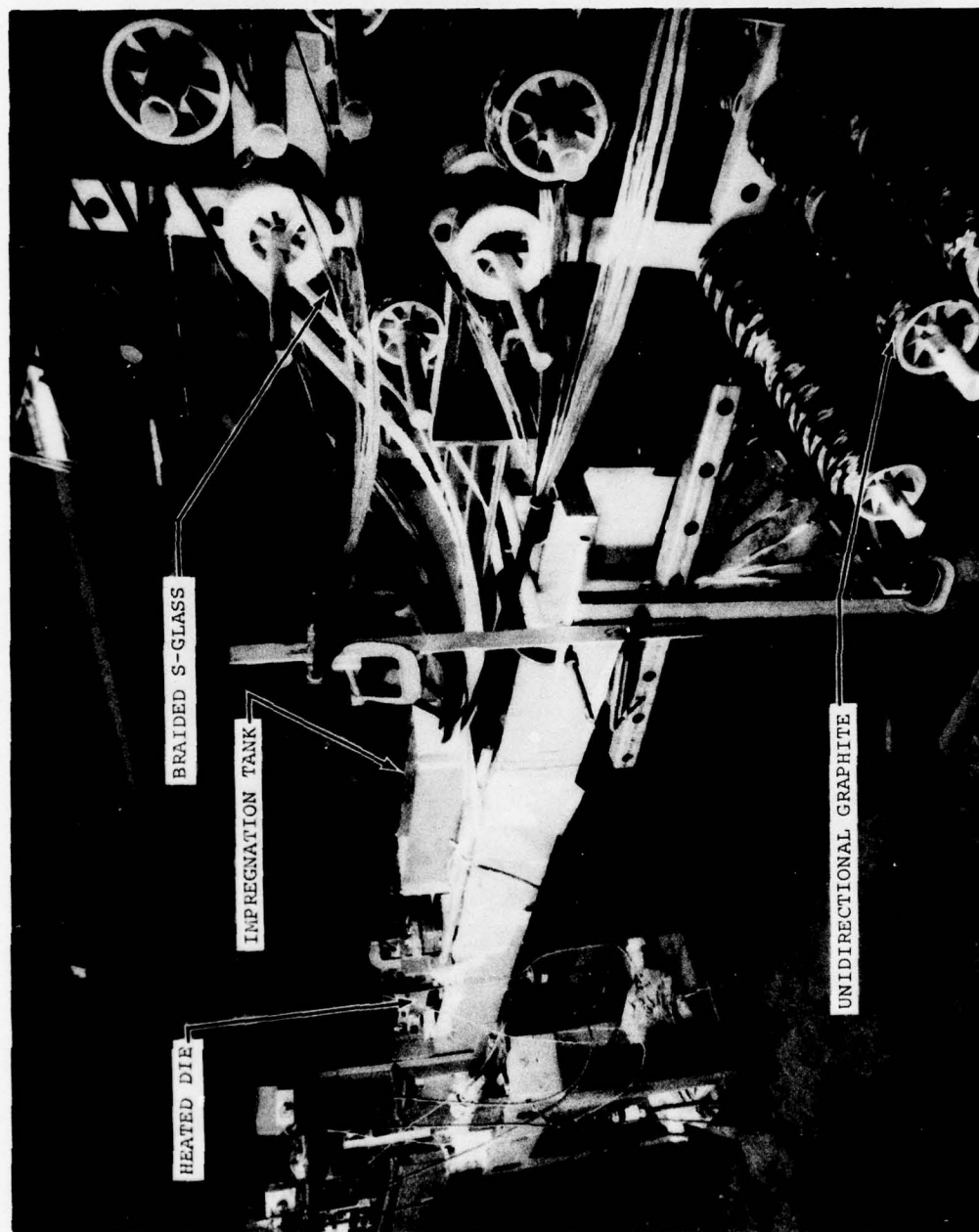
Figure 35. Pultrusion of Demonstration Article
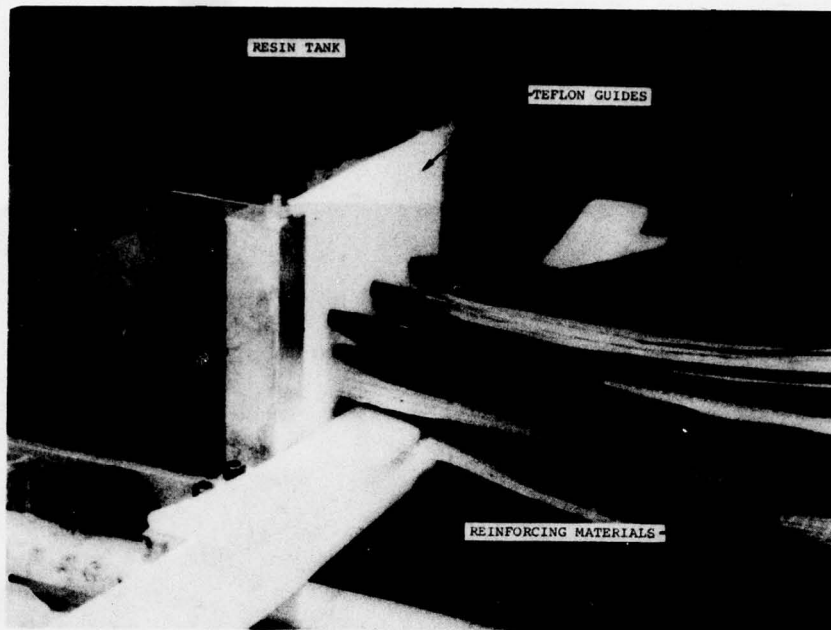
Figure 36. Entrance to Resin Tank.
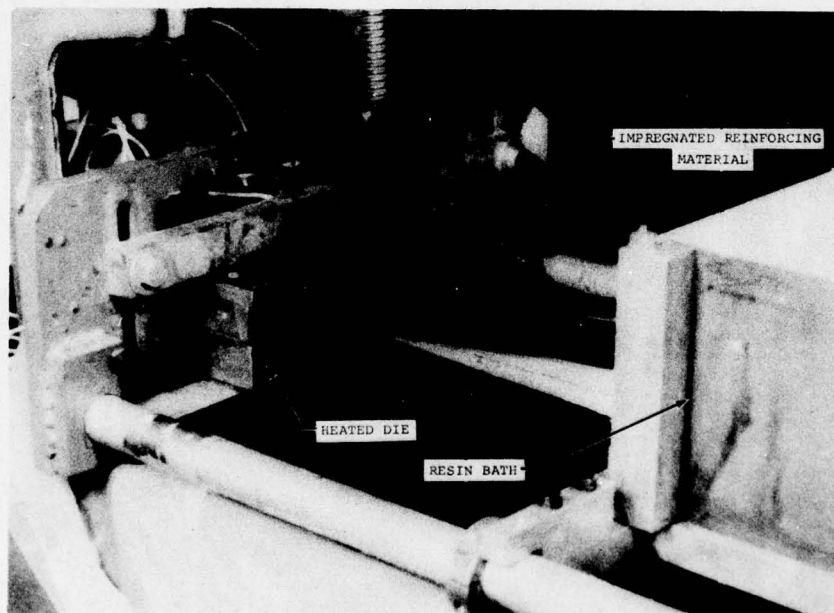
Figure 37. Entrance to Pultrusion Die.

74
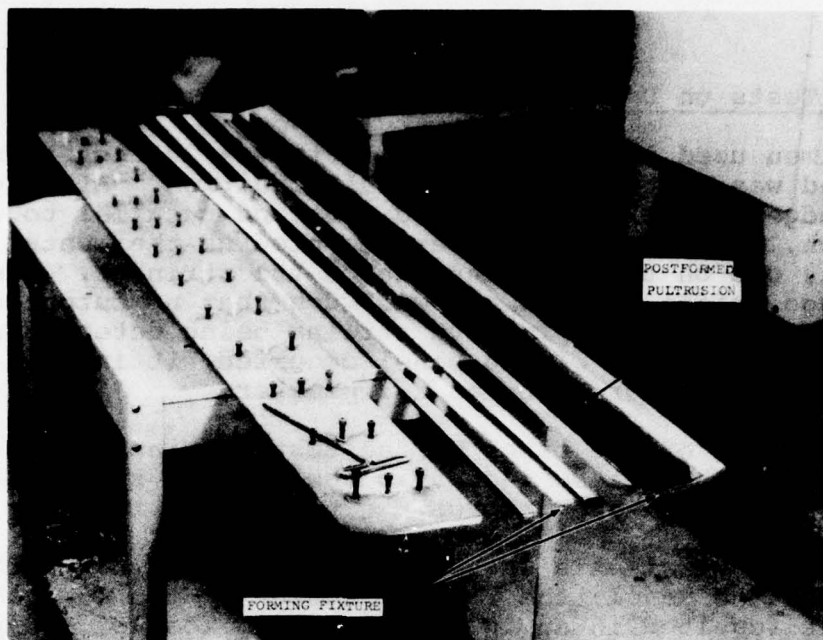
Figure 38. Postforming Tool.
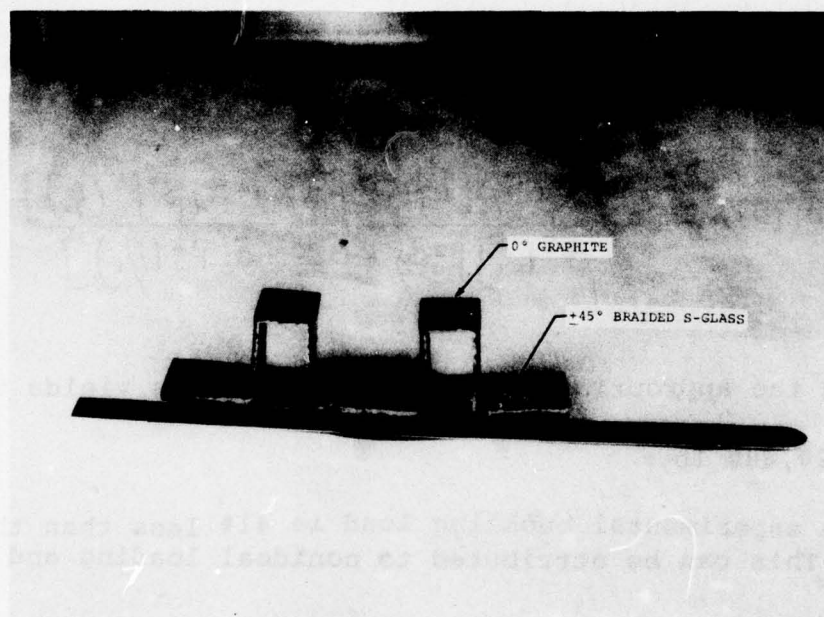
Figure 39. Section of Postformed Material.

75

Buckling Tests on Demonstration Article

The specimen used in the buckling tests was 20 inches in length and was postcured in the flat condition. After grinding the edges (short) to be loaded flat and parallel to within 0.005-inch, two strain gages were mounted on the center of the panel, one on the skin and one on the stringer. The strain gage bridge arrangement was such that no output occurred provided the compressive strains being detected were equal. When buckling of the panel occurred, it induced a signal that was recorded on an X-Y plotter.

The arrangement of the panel in the test machine may be seen in Figure 40, which also indicates the loading system (based upon pins acting on vee grooves).

The elastic neutral axis was calculated to be 0.239-inch from the face of the panel; the line of action of the load through this point resulted in an initial buckling load of 700 pounds and an instability load of 9950 pounds. Adjusting the load application points to a distance of 0.225-inch from the panel face resulted in the buckling load occurring at 4100 pounds and the instability load at 12,340 pounds. These results are shown in the load-deflection trace of Figure 41. P critical can also be calculated from Equation (C-30) presented in Appendix C:

$$P_{cr} = \sqrt{D_{11} D_{22}} \; \frac{{}_0\int^b \phi \, dy + b \sum_{i=1}^{N} \left[j_i \beta^4 f^2(\eta_i) + x_i \beta^2 f'^2(\eta_i) \right]}{\beta^2 \left[{}_0\int^b f^2 dy + b \sum_{i=1}^{N} \delta_i f^2(\eta_i) \right]}$$

Inserting the appropriate structural properties yields the result,

$$P_{cr} = 20,880 \text{ lb.}$$

Thus, the experimental buckling load is 41% less than that predicted. This can be attributed to nonideal loading and efge effects.

Crippling Tests on Demonstration Article

Small angle and channel coupons were cut from the final part and tested in compression for crippling strength; the results are presented in Table 16 and plotted as dimensionless ratios
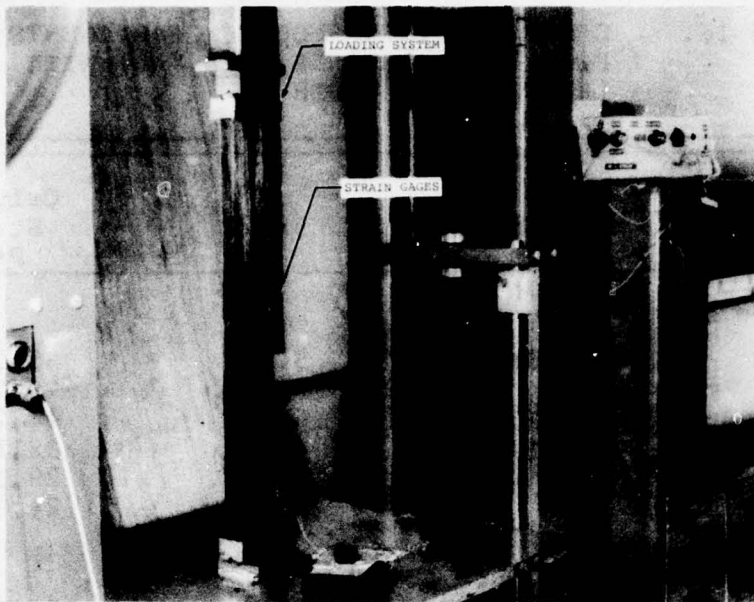
Figure 40.   Panel Undergoing Buckling Tests.
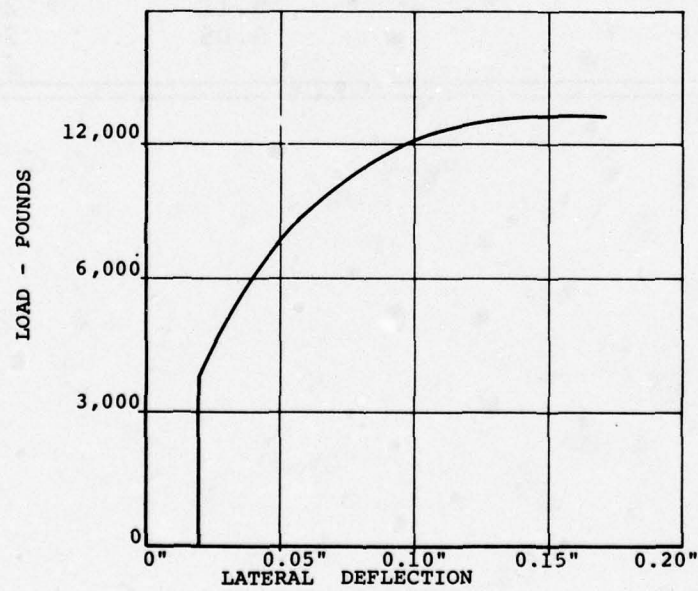
Figure 41.   Load Deflection Relationships
Obtained During Buckling Tests.

TABLE 16.	CRIPPLING TEST RESULTS OF ELEMENTS FROM FINAL ARTICLE - $(0, \pm 45)$ WITH LENGTH = 4 x WIDTH

Test No.	Shape	b/t	Crippling Stress Psi
One Edge Free			
155-1	Angle	6.66	35,390
155-2	"	6.94	28,710
155-3	"	6.54	44,210
155-4	"	4.16	32,160
155-5	"	3.84	29,180
155-6	"	4.31	24,330
No Edge Free			
155-7	Channel	7.34	21,120
155-8	"	7.08	32,200
155-9	"	7.46	22,620
155-10	"	9.13	27,000
155-11	"	9.11	26,770
155-12	"	9.11	26,200
155-13	"	9.05	24,100

Material	E_0^C (10^6psi)	E_{90}^C (10^6psi)	G (10^6psi)	ν_{12}	ε_0^{cu} $(\mu\text{in/in})$	ρ (lb/in^3)
Modmar II/5206	23	1.4	.7	.24	11,900	.055
Hercules AS/3002	16	3.7	.8	.25	13,500	.055
Celanese GY-70/904	42	.9	.6	.25	2,100	.061
120 Fiber-glass cloth/ MIL-R-9300 Epoxy	2.2	2.2	.6	.11	20,000	.065

TABLE 17. UNIDIRECTIONAL PROPERTIES OF COMPOSITE MATERIALS USED IN EFFICIENCY ANALYSIS

in Figure 42. Due to the limitations imposed by the available specimen size, the b/t ratios are considered to be too low to demonstrate any significant changes in the relationship between the two parameters. The results are generally somewhat lower than that expected.
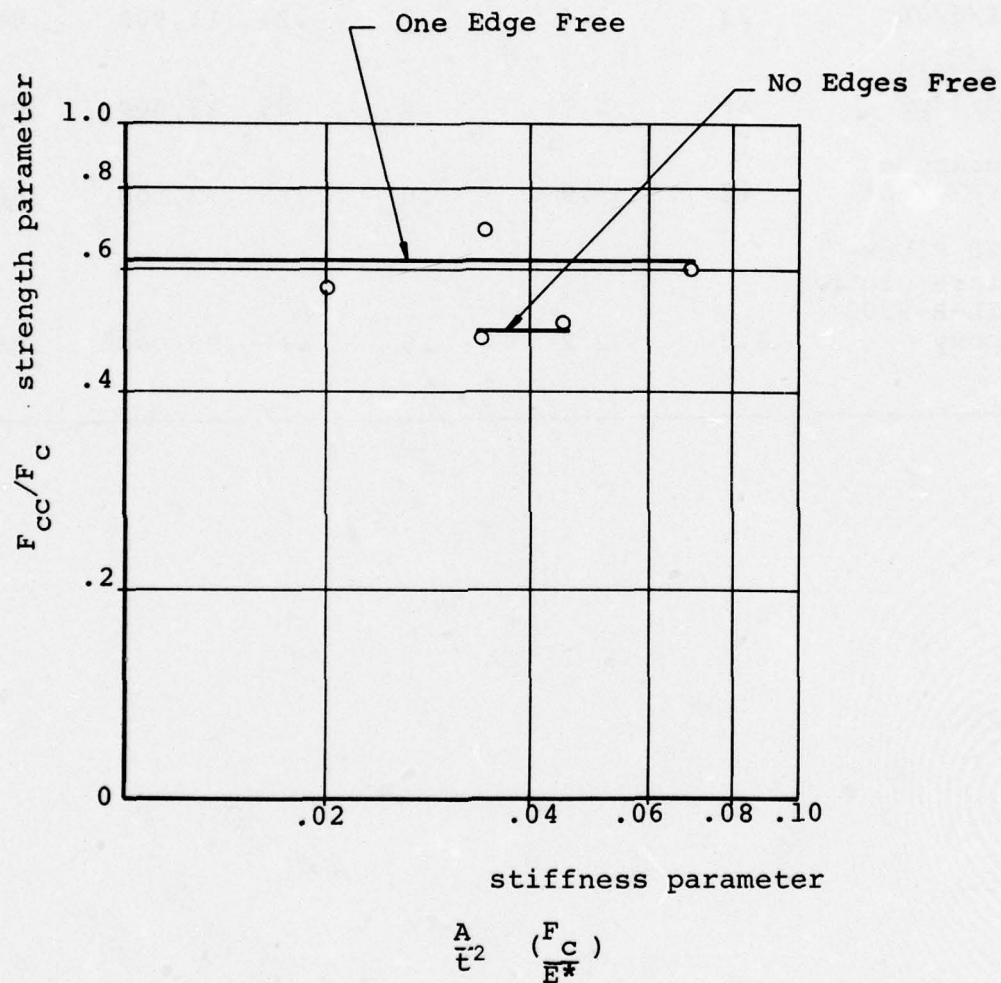
Figure 42. Crippling Stress of (0, ±45) Hybrid Composite.

APPLICATION STUDIES

The application study established the potential performance
and cost advantages that the pultrusion process would offer in
manufacturing high-performance aircraft structural components.
The approach was to parametrically evaluate composites for
specific structural applications and then to develop conceptual
designs based on the parametric studies and manufacturing lim-
itations of pultrusion.

PARAMETRIC STUDY

To assess the potential of various fibers, matrices and
structural proportions, a structural efficiency analysis
approach was used. The low density and high strength and
stiffness of composites, relative to conventional aerospace
metals, indicates that they should be highly efficient as
structural members. For a unidirectional intermediate
strength graphite/epoxy, the strength-to-density ratio in the
direction of the fibers is in the order of 3.6×10^6 inches.
This value compares to 0.7×10^6 inches for 7075-T6 aluminum
alloy. The conclusion drawn from the values that a graphite/
epoxy tension member would be 1/7 the weight of an aluminum
member would most likely be invalid since other design con-
siderations such as secondary loads, minimum gage, eccentric-
ity of load, etc., would completely change the relationships.

Various techniques have been used to determine optimum designs
for structural elements constructed from conventional isotrop-
ic metals. These techniques, for many reasons, are insuffi-
cient for the optimum design of composite structural elements.
Unlike conventional metals, filamentary reinforced composite
materials are orthotropic, or even anisotropic. The full
determination of the appropriate form and optimum proportions
of structural elements using these materials requires consid-
eration of significantly larger numbers of design variables
than do isotropic designs. Imposing undue restrictions on the
design process just to effect a simpler solution to the
optimization problem excludes many of the advantages offered
by using composite materials.

Integrally Stiffened Panels

The structural efficiency analysis used involved the deter-
mination of generalized weights of the integrally stiffened
panels required to carry given axial loading intensities.

The appropriate parameters for this generalization have been
found to be weight per unit surface area divided by panel
width as a function of axial running load per unit width.
Evaluation of the minimum weight form and proportions in each
case required the application of appropriate failure criteria.
Overall panel stability was described by orthotropic stiffened
plate theory; local stability of the panel elements was de-
scribed by orthotropic plate theory.[15] A maximum strain
criterion was used to describe strength limitations of the
composite materials.

The optimum minimum weight configuration is obtained when two
failure modes occur simultaneously for a given load. An ex-
ample would be the panel elements failing by local instability
at the same stress level that overall buckling occurs. The
solutions are obtained graphically by cross plotting the
weight parameter and the load parameter in terms of panel
dimension ratios.

Two panel configurations were studied, a webcore panel and a
single-face corrugated panel, as defined in Figures 43 and 44.

To reduce the number of variables, certain of the panel
dimensions were either held constant or related to another
variable. For the webcore panel, the skin and webs were the
same thickness. The height of the web was related to the web
spacing such that the web and skin elements had the same
resistance to local buckling. The panel width-to-height ratio
was maintained at $b/a = 2$, representative of a panel element
between frames in a fuselage. For the single-face corrugated
panel, the height and width of the stiffener were maintained
equal. The space between stiffeners was related to the
stiffener height such that both elements had the same resis-
tance to local buckling. These restrictions on dimensions
minimize the weight as regards local buckling by ensuring
that all material fails at the same time.

The load parameter for overall stability of the webcore panel
is

$$\frac{N_x}{b} = \frac{\pi^2 E_x}{2\,m^2}\left(\frac{1 + \sqrt[4]{d}}{1 - \nu_{xy}\nu_{yx}}\right)\left(\frac{t}{d}\right)\left(\frac{d}{a}\right)^3\left(\frac{a}{b}\right)^5 (A+B) \qquad (1)$$

where

A and B are defined in Appendix C.

82
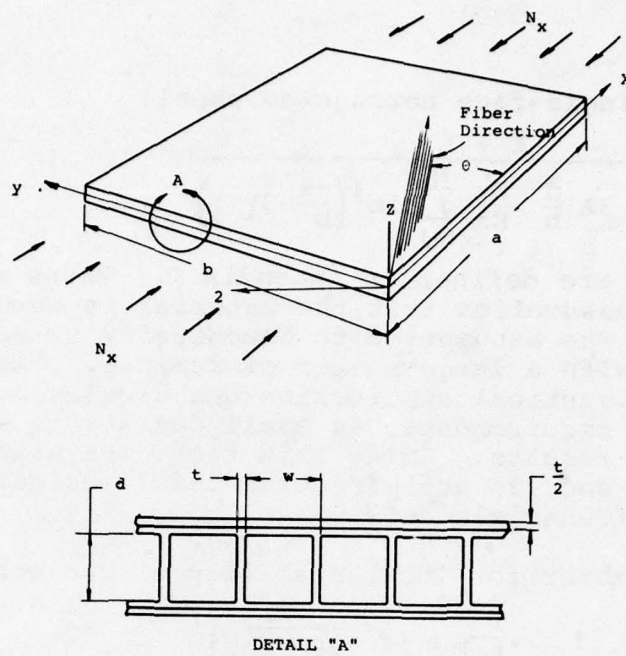
Figure 43. Webcore Panel - Dimensions and Coordinate System.
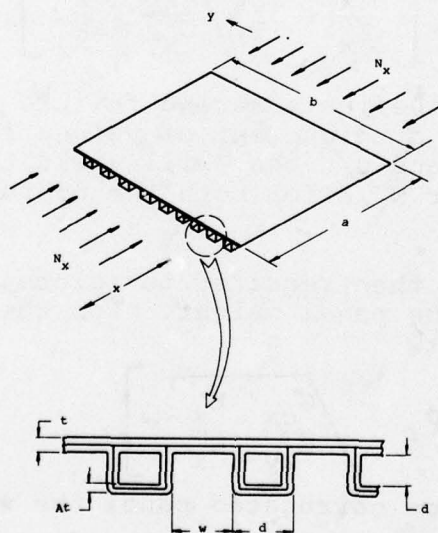
Figure 44. Single-Face Corrugated Skin Panel - Dimensions and Coordinate System.

83

and for the single-face corrugated panel:

$$\frac{Nx}{b} = \frac{K + L}{\frac{1}{2} + 3A \frac{d}{b} \frac{E'_x}{E_x} \sum_{i=1}^{N} \sin^2\left(\frac{\eta\pi}{b} \eta_i\right)}$$ (2)

where K and L are defined in Appendix C. These equations are based on the assumption that the material is homogeneous and orthotropic. The assumption to homogeneity is met for a balanced lay-up with a large number of laminae. The sequence of laminae in a practical application can usually be arranged to satisfy these requirements, as small deviations will not grossly affect the results. Using this fact, the elastic moduli in equations (1) and (2) are effective moduli calculated for the plate and stiffener elements.

The load parameter for local stability of the webcore panel is

$$\frac{N_x}{b} = \left(\frac{t}{d}\right)^3 \left(\frac{d}{b}\right) G'_{cx} \left[2\frac{E_x}{E'_x} + \sqrt{\frac{G'_{cx}E_x}{G_{xy}E'_x}}\right]$$ (3)

and for the single face corrugated panel is

$$\frac{N_x}{b} = \left(\frac{t}{d}\right)^3 \frac{d}{b} G'_{xy} \left[A^2\frac{E_x}{E'_x} + 3A^4 \sqrt{\frac{G'_{xy}}{G_{xy}} \frac{E_x}{E'_x}}\right]$$ (4)

Equations (3) and (4) define a second failure criterion for the panels. There is a unique set of values for the panel dimension ratios b/d and d/t that will yield the same value for the load parameter N_x/b for both the overall and local instability condition.

A weight parameter is then required to relate the panel dimension ratios to the panel weight. For the webcore panel the weight parameter is

$$\frac{W}{ab^2} = \frac{t}{d} \frac{d}{b} \left[2 \rho_f + \rho_r \sqrt{\frac{G'_{cx}}{G_{xy}} \frac{E_x}{E'_x}}\right]$$ (5)

and for the single-face corrugated panel the weight parameter is

$$\frac{W}{ab^2} = \frac{t}{d} \; \frac{d}{b} \; \left[\varrho_p + 3A^2 \varrho_s \; \sqrt{\frac{G'_{xy}}{G_{xy}} \; \frac{E_x}{E'_x}} \right] \tag{6}$$

By solving Equations (2), (4) and (6) for various b/d and d/t ratios and substituting in the elastic properties of the composite, a family of curves is obtained that relates panel weight and dimensions to the ability to transmit an axial compressive load. In Figure 45 the results are plotted for a single-face corrugated panel based on the properties of Hercules AS graphite fiber and epoxy resin. Half the fibers in the plate and stringers are oriented 0° to the load and the remaining half are oriented $\pm$ 45° to the load. For clarity, only the pertinent curves are plotted. Three points of common intersection can be noted for different sets of dimension ratios and parameters, as follows:

1. $\dfrac{N_x}{b^2} = 280$ for $\dfrac{b}{d} = 20$, $\dfrac{d}{t} = 3$

2. $\dfrac{N_x}{b^2} = 24$ for $\dfrac{b}{d} = 40$, $\dfrac{d}{t} = 6$

3. $\dfrac{N_x}{b^2} = 5.7$ for $\dfrac{b}{d} = 60$, $\dfrac{d}{t} = 9$

A line drawn through these points defines the minimum weight panel that will carry any given load.

A third failure criterion of ultimate compressive strength can also be applied. The load parameter for a webcore panel is

$$\frac{N_x}{b} = e^{cu}_{0^\circ} \left(\frac{t}{d}\right) \left(\frac{d}{b}\right) \left[2E_x + E'_x \sqrt{\frac{G'_{cx}}{G_{xy}} \; \frac{E_x}{E'_x}} \right] \tag{7}$$

and for a single-face corrugated panel it is

85
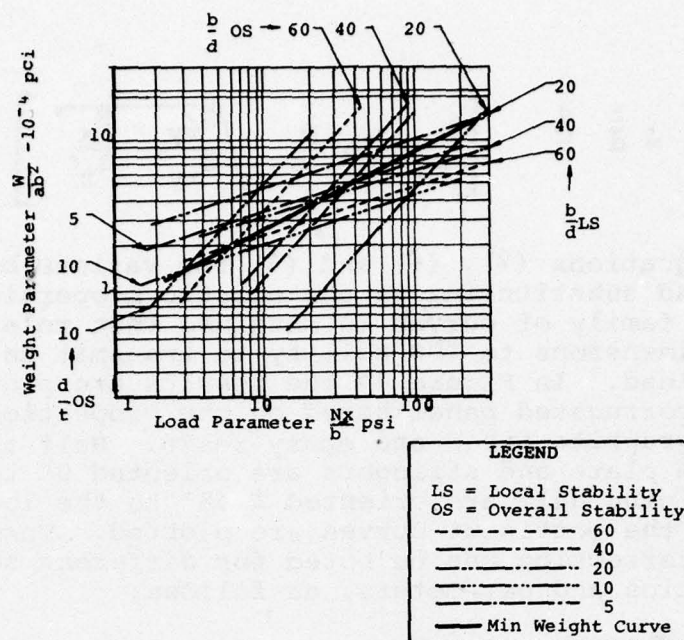
Figure 45.   Minimum Weight Curves for Structural
Efficiency Analysis.
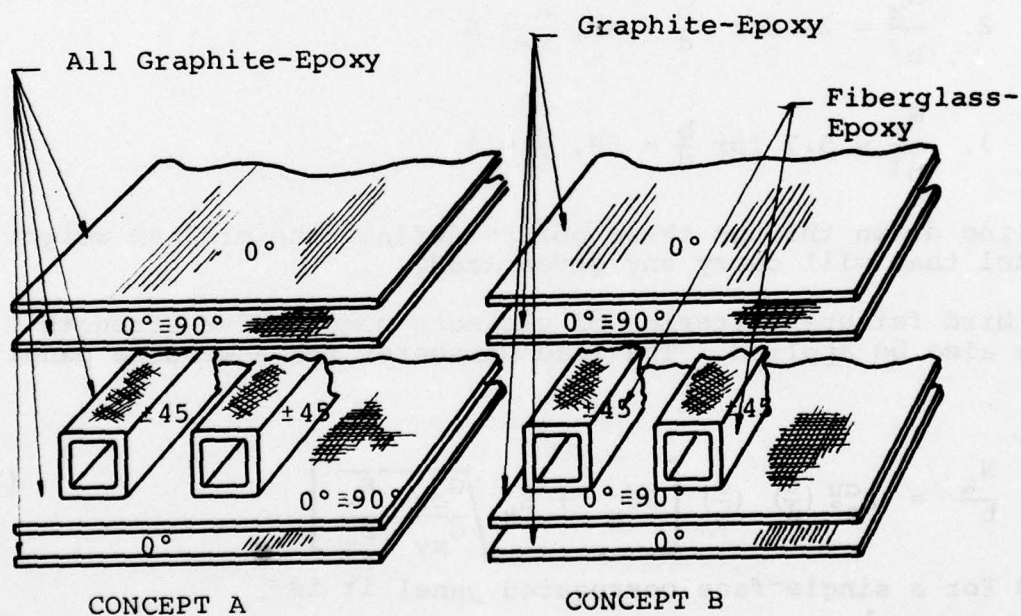
Figure 46.   Webcore Designs Analyzed.

$$\frac{N_x}{b} = \frac{t}{d} \frac{d}{b} \varepsilon_{0°}^{cu} \left[E_x + 3E_x' A^2 \sqrt{\frac{G_{xy}'}{G_{xy}} \frac{E_x}{E_x'}} \right] \quad (8)$$

The ultimate strength criterion interacts with the overall stability to set the minimum weight curve since, as the elements thicken, it precludes local instability.

Practical designs must also consider the minimum gage that can be used for a material in a particular application. The minimum gage is set by secondary factors such as manufacturing requirements, ballistic damage tolerance, foreign object impacts and resistance to environment to name a few. Since minimum gage cannot be considered in the context of the loads, it is included in the structural efficiency analysis when considering a specific application. The effect of minimum gage was included by considering a helicopter fuselage application. In studies of the application of composite materials to helicopter fuselages by Sikorsky Aircraft[16] and Boeing Vertol,[17] the frame spacing in both cases was 20 inches. By assuming a = 20 inches and considering two minimum gages as t = .020 inch and t = .040 inch, a lower bound is defined for the minimum weight curve.

The mechanical properties used in this study are shown in Table 17. Using laminated plate theory, the elastic constants for the composite material were calculated. Then, by assuming the plate to be orthotropic and homogeneous, the equivalent elastic modulii were calculated.

For a webcore panel, two concepts were studied as shown in Figure 46. The graphite-epoxy properties used were for Modmar II/5206 and the fiberglass-epoxy was with E-glass 120 cloth. The results of the calculations are presented in Figure 47. The effects due to local stability are cut off by the minimum gage lines, and the sloped portion for $N_x/b > 80$ is set by ultimate compressive strength and overall stability. Superimposed are the load and weight ranges reported[16] for the CH-53D helicopter fuselage converted to efficiency parameters. For this application, the composite webcore skin at 0.020-inch thick would show a 19% weight savings over the areas with 0.032-inch aluminum skin and be equal for the areas with 0.025-inch skin. However, a minimum gage of 0.020-inch would
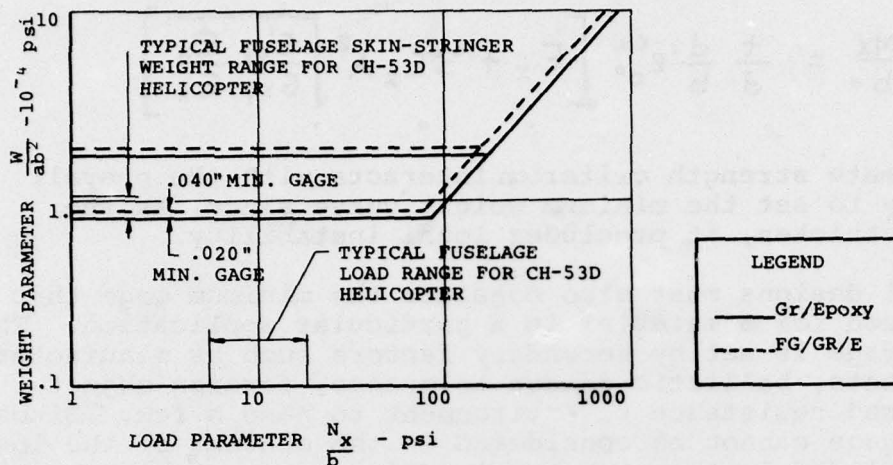
Figure 47. Structural Weight Relation
for Webcore Panel.
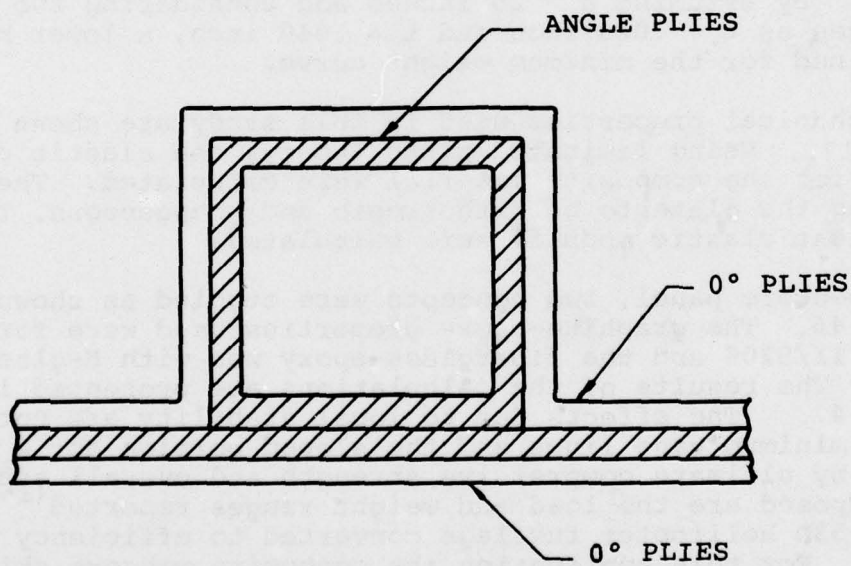
Figure 48. Distribution of Plies in Single-Face
Corrugated Panels.

probably not be satisfactory when all the design considerations were included. It is of interest to note the small weight penalty incurred by using fiberglass for the webs and part of the skin thickness, pointing out the viability of the hybrid fiber approach.

The webcore panels will be effective at higher loadings than would be found on wing and tail surface skins.

The number of cases were run for single-face corrugated panels to explore their weight sensitivity to various parameters. The ply distribution within the laminate is shown in Figure 48. The actual ratio of 0° plies to angle plies was varied. In the first case, the stringer thickness to plate thickness ratio was 0.5 and the angle plies represented 50 percent of the respective thicknesses. The results are plotted in Figure 49. The use of fiberglass for the angle plies in this case causes a 50-percent increase in structural weight over the all graphite panel. The graphite panel is 40 percent heavier than the aluminum skin-stringer panel.

Next, the ratio of $0°:\pm 45°$ plies was changed from 1:1 to 2:1, thereby increasing the modulus in the load direction. The results are plotted in Figure 50. The solid line is the same as that in Figure 49. In this case the weight of the graphite-epoxy composite panel has been reduced and now is only 15 percent heavier than the aluminum skin-stringer panel. In the balance of the cases studied, the thickness ratio for $0°:\pm 45°$ was maintained at 2:1.

The modulus of the fiber should have a marked effect on the response of structures to buckling. Therefore, the efficiency was calculated for a low, medium and high modulus graphite. The material properties used were for Hercules AS/3002, Modmar II/5206 and Celanese GY-70/904 epoxy systems. The results are plotted in Figure 51. The graphite-epoxy composite panels based on Modmar II and Hercules AS fibers had essentially the same efficiency. However, the Celanese GY-70 fiber-based composite showed a marked reduction in weight of 35 percent. This panel would average about 10 percent lighter than the aluminum skin-stringer panels for the CH-53D helicopter.

The orientation of the angle plies has a marked effect on buckling. Four different fiber orientations were compared to evaluate this effect. The orientations chosen were $\pm 45°$, $\pm 60°$, $\pm 30°$ and $0°,90°$. The results of the calculations are shown in Figure 52 for a Hercules AS/904 composite. The $\pm 45°$ orientation for the angle plies results in the lowest weight for a given load. This result is primarily influenced by the

89
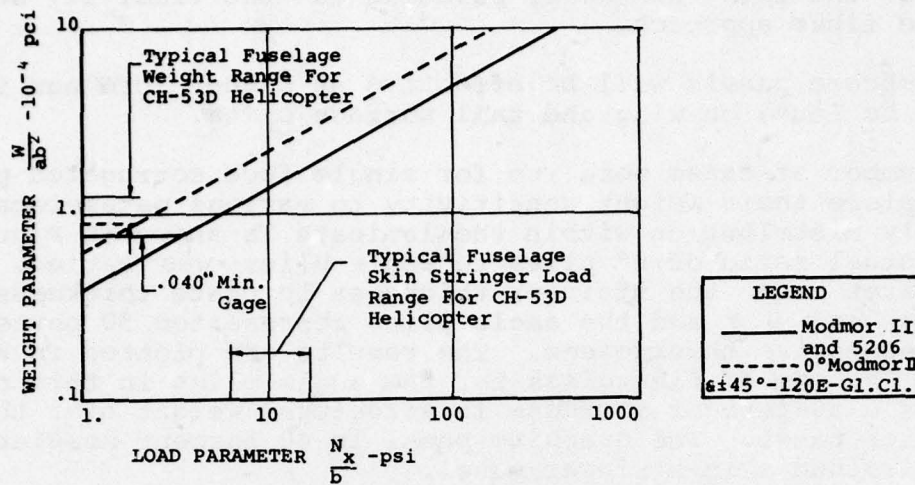
Figure 49. Fiberglass Vs. Graphite for Angle Ply Layers With Single-Face Corrugated Panels.
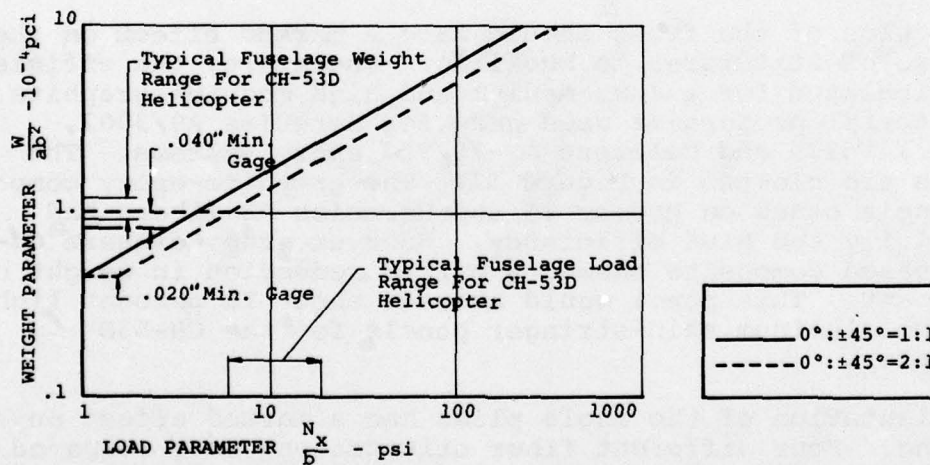
Figure 50.   Thickness Effect in Single-Face Corrugated Panels.
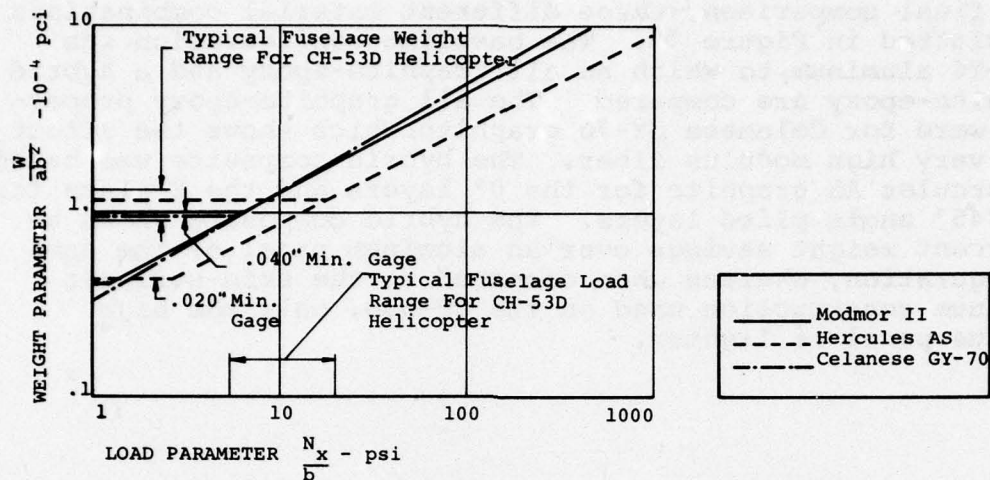
Figure 51. Modulus Effect in Single-Face
Corrugated Panels.
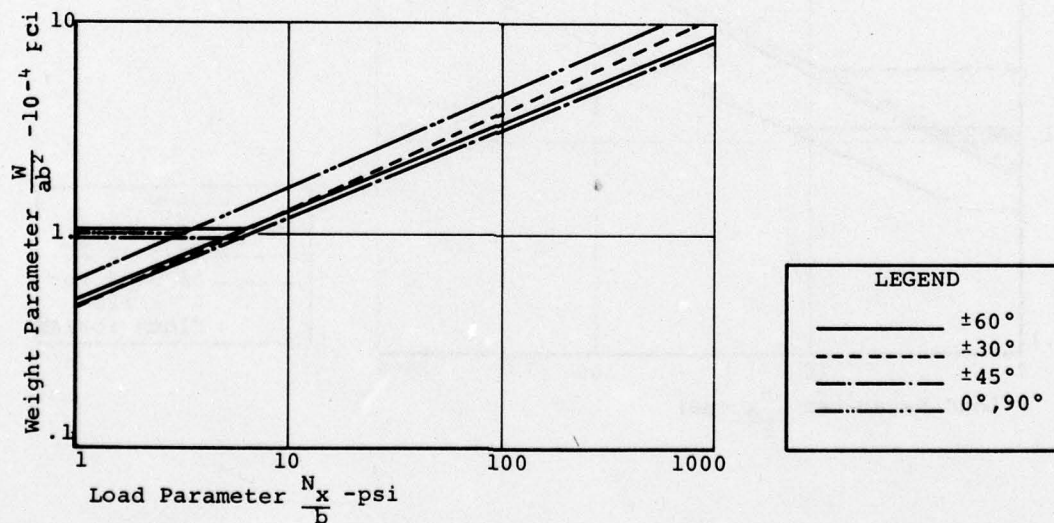
Figure 52. Orientation Effect of Fibers for
Single-Face Corrugated Panels.

91

Load Parameter for local buckling, which is directly dependent on the shear modulus of the material.

As a final comparison, three different material combinations are plotted in Figure 53. The baseline configuration was 7075-T6 aluminum, to which an all graphite-epoxy and a hybrid graphite-epoxy are compared. The all graphite-epoxy properties were for Celanese GY-70 graphite, which shows the effect of a very high modulus fiber. The hybrid composite was based on Hercules AS graphite for the 0° layers and the E-glass for the ±45° angle plied layers. The hybrid composite shows an 8-percent weight savings over an aluminum panel of the same configuration, whereas when compared to the skin-stringer aluminum construction used on the CH-53D, only the high modulus panel is lighter.
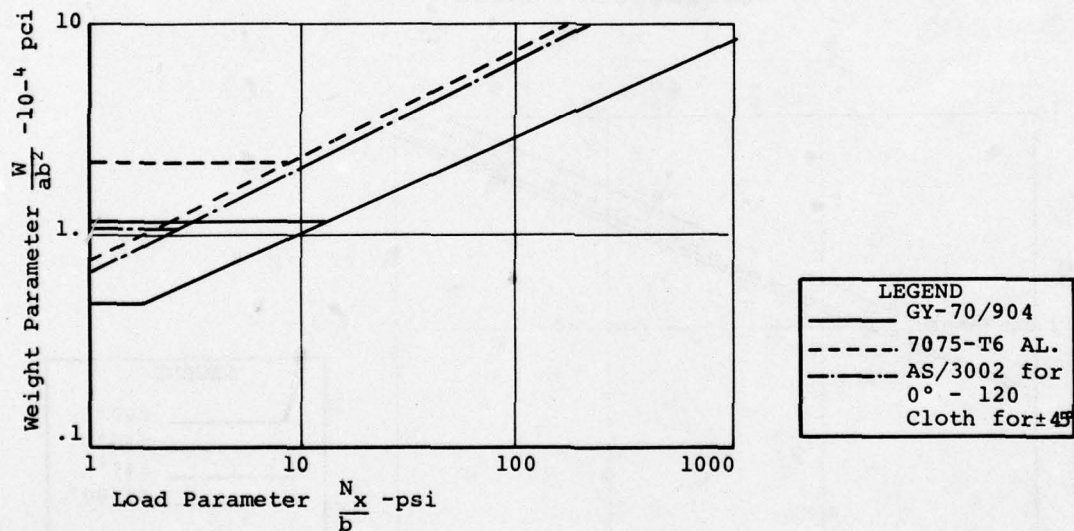
Figure 53. Material Comparison for Single-
Face Corrugated Panels.
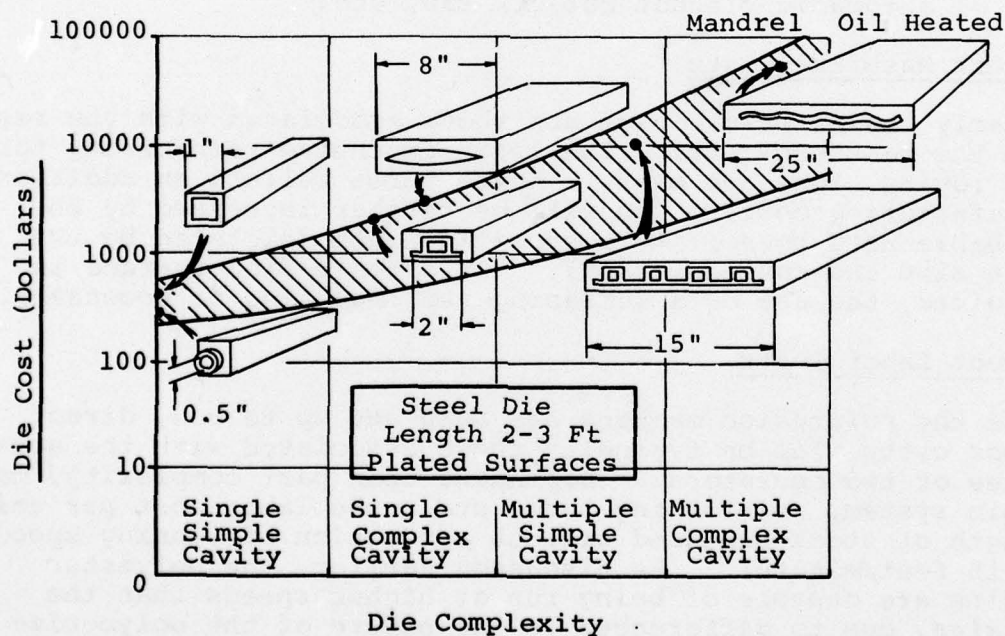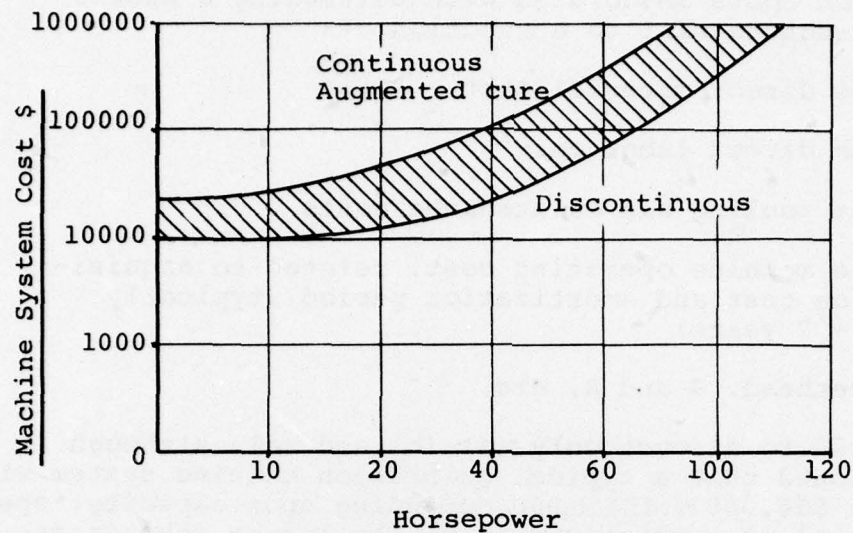
Figure 54. Characteristic Die Costs.

MANUFACTURING COST STUDIES

The production costs associated with pultruding a specil
length of structure will be a function of:

    (a)   the direct material cost

    (b)   the direct labor cost

    (c)   the tooling and maintenance costs

    (d)   the machine operating cost, related to acquisi-
          tion cost and amortization period (typically
          5 - 7 years)

    (e)   overhead, G and A, etc.

It is proposed to discuss only (a), (b) and (c), although it
may be mentioned that a typical pultrusion machine system will
cost between $30,000 and $150,000, depending upon capacity, speed,
power and level of sophistication. The latter item is re-
lated to the use of continuous or discontinuous pullers, use
of a preheating system (dielectric, inductive, infrared, etc.),
use of automatic product cut-off saw, etc.

Direct Material Costs

Clearly the material costs are those associated with the resin
and the reinforcement. The least expensive reinforcing forms
are roving, tows and mats. Woven forms reflect an additional
manufacturing cost, which will be further increased by the
probable need to slit to a specific width (dictated by the
part size and configuration). If a resin-rich surface is
required, the use of a surfacing veil may also be necessary.

Direct Labor Costs

Once the pultrusion machine has been set up to run, direct
labor costs will be typically those associated with the ser-
vices of two operators. Dependent upon part complexity, mass,
resin system, reinforcing form, etc., the labor cost per unit
length of stock produced will be a function of running speed
(1-15 feet/minute). As discussed earlier, the polyester
resins are capable of being run at higher speeds than the
epoxies, due to differences in the nature of the polymeriz-
ation chemistry for these materials. Products involving
woven forms generally are run at lower speeds, due to con-
straints on material feed-in and the need for adequate resin
impregnation.

Tooling and Maintenance Costs

The costs of tooling are those due to dies, special heaters, mandrels (for hollow products), creels, in-feed devices (for reinforcing materials), special resin impregnation systems (tanks, pressurizing units), shaped gripper faces, etc.

With the exception of the die, most production requirements can be met with standard equipment readily adapted to meet changing needs. Die costs are highly variable, depending upon complexity and size. As a generality, such costs may vary from $1,000 to $50,000. To illustrate this concept, Figure 54 shows the acquisition costs of typical dies purchased recently, related to geometric and size factors.

The maintenance costs of dies are generally related to such needs as cleaning and replating and will not normally exceed 10% of acquisition cost per 100,000 feet of stock produced.

CONCLUSIONS

This program enables the following conclusions to be reached:

1. The use of existing epoxy resin systems, of the epichlorohydrin bisphenol A type, are suitable for use in the pultrusion process. In order to minimize die adherence, the use of diamine hardeners appears beneficial.

2. From the experience obtained during this program, the use of internal mold releases is not effective and will degrade mechanical properties.

3. The use of polyester resins with graphite reinforcements should include pretreatment of the graphite to enhance filament-matrix bonding.

4. The viscosity of the resin system for optimum processing should be in the range of 300-1000 cps, depending upon product size and form of reinforcing.

5. When pultruding with the epoxy resin systems investigated during this contract, stock speeds of 1 to 2 ft/min would produce a level of cure such that the material would be totally solid and tack-free. Fully-cured conditions could be attained by simply subjecting the product to a normal post-curing cycle.

6. The mechanical properties exhibited by the specimens produced by pultrusion were generally superior or equal to those exhibited by equivalent specimens produced by vacuum bagging.

7. While experience with the resin systems intended for prepregging demonstrated some difficulties in processing, the mechanical properties exhibited by these materials were very satisfactory. This suggests that reformulation of such a resin system for use in pultrusion may be advantageous.

8. For the epoxy resin systems investigated, post-forming could easily be carried out during the final curing cycle, provided no curvature was induced along the axes of the filaments.

9. The static and fatigue properties demonstrated by the specimens pultruded were consistent with those

characteristic of other processing techniques.
It may, therefore, be asserted that pultrusion
does not intrinsically degrade the short- or long-
term behavior of a composite material.

10. In the absence of ply-on-ply biased reinforcement,
the use of braiding to provide a convenient form for
pultrusion is viewed as acceptable.

11. Complex, hybrid structures may be produced by pul-
trusion, which is both cost-effective and efficient.
Any limitations inherent in the process, from a ma-
terial orientation standpoint, are offset by the
cost and repeatability of the technique.   Good
translation of coupon properties into larger scale
structures appears readily attainable.

RECOMMENDATIONS

Based on the information derived from this program, it is recommended that:

1. Effort be directed toward the formulation of high-performance resins specifically suited to pultrusion processing requirements, which possess good processability and mechanical properties at low and elevated temperatures.

2. Advanced design studies be initiated to enable pultruded substructures to be incorporated into major assemblies.

3. Programs be instituted to develop methods for producing and in-feeding multi-oriented reinforcements to remove the existing limitation on available reinforcing forms.

98

REFERENCES CITED

1. Dhonau, O.E., CONCEPTUAL DESIGN OF COMPOSITE V/STOL TACTICAL FIGHTER, LTV Corporation, AFML TR 73-3, Air Force Material Laboratory, Wright-Patterson Air Force Base, Ohio, July 1973.

2. Ewald, G.W., PULTRUSION, MODERN PLASTICS ENCYCLOPEDIA 75-76, Vol. 52, October 1975.

3. Goldsworthy, W.B., PULTRUSION - A GROWING TONNAGE MARKET FOR REINFORCED PLASTICS END PRODUCTS, Reinforced Plastics (London), June 1971.

4. Jones, B.H., PULTRUDING FILAMENTARY COMPOSITES, AN EXPERIMENTAL AND ANALYTICAL DETERMINATION OF PROCESS PARAMETERS, SPI 29th Annual Technical Conference, Reinforced Plastics Composites Institute, February 1974.

5. Maximovich, M.G., Hollingsworth, C.D., Neff, R.M., DEVELOPMENT OF IMPROVED ENVIRONMENTAL RESISTANT ORGANIC-REINFORCED MATRIX SYSTEMS, SPE Pactec 75, Las Vegas, 1975.

6. Oleesky, S., and Mohr, G., HANDBOOK OF REINFORCED PLASTICS OF THE SPI, Reinhold Publishing Co., 1964.

7. Timoshenko, S., and Goodier, J.N., THEORY OF ELASTICITY, McGraw Hill Book Company, 1951.

8. Anon., ADVANCED COMPOSITES DESIGN GUIDE (3rd Ed.), Air Force Materials Laboratory, Vol. IV, 1973.

9. Boller, K.H., SOME FATIGUE CHARACTERISTICS OF GLASS REINFORCED COMPOSITE LAMINATES, ASME Design Engineering Conference, Pages 70-DE-6, 1970.

10. Rao, N. and Hofer, K.E., FATIGUE BEHAVIOR OF GRAPHITE/ GLASS/EPOXY COMPOSITES, Final Report on Contract No. N00019-72-C-0294, Naval Air Systems Command, April 1973.

11. Anon., PLASTICS FOR FLIGHT VEHICLES PART I REINFORCED PLASTICS, MIL-HDBK-17, 1959.

12. Novak, R.C., TORSIONAL FATIGUE BEHAVIOR OF UNIDIRECTIONAL RESIN MATRIX COMPOSITES, Composite Materials Testing and Design, ASTM, STP 546, 1974.

13. Anon., METALLIC MATERIALS AND ELEMENTS FOR AEROSPACE VEHICLE STRUCTURE, MIL-HDBK-5B, 1972,

14. Jones, B.H., PROBABILISTIC DESIGN AND RELIABILITY, in Composite Materials, Vol. 8, Structural Design and Analysis Part II, Academic Press, New York, 1975.

15. Lekhnitskii, S.C., Anisotropic Plates, Gordon and Breach, 1968.

16. Rich, M.J., et al, APPLICATION OF COMPOSITES TO HELI-COPTER AIRFRAMES AND LANDING GEAR STRUCTURES, Sikorsky Air-Craft, NASA CR-112333, National Aeronautical and Space Administration, Langley, Virginia, June 1973.

17. Swatton, S., STUDY OF ADVANCED STRUCTURAL CONCEPTS FOR FUSELAGE, Final Report on Contract DAA-JO2-72-C-0056, U.S. Army Air Mobility Research and Development Laboratory, October 1973.

18. Cagle, C.V.(Ed.), HANDBOOK OF ADHESIVE BONDING, McGraw Hill Book Co., 1973.

19. Timoshenko, S.P., and Gere, J.M., Theory of Elastic Stability, McGraw Hill Book Co., 1961.

20. Roark, R.J., Formulas for Stress and Strain, McGraw-Hill Book Co., 1954.

APPENDIX A

CHEMICAL CONSIDERATIONS FOR
EPOXY RESINS WHEN BEING USED IN PULTRUSION

Since 1950, the epoxy resins have become increasingly important to the reinforced plastics industry. Although many different types of epoxy resins can be produced, three are of major commercial importance: (1) conventional epoxy resins, based on epichlorohydrin and bisphenol A; (2) epoxy novolac resins; and (3) cycloaliphatic epoxy resins. The conventional resins are generally the most commonly used, and are now the least expensive to manufacture. This is presently the most suitable epoxy system for use in pultrusion; some of its properties in relation to the process are examined below:

The conventional epoxy resin is synthesized by adding epichlorohydrin to bisphenol A in the presence of sodium hydroxide to remove the hydrochloric acid given off.
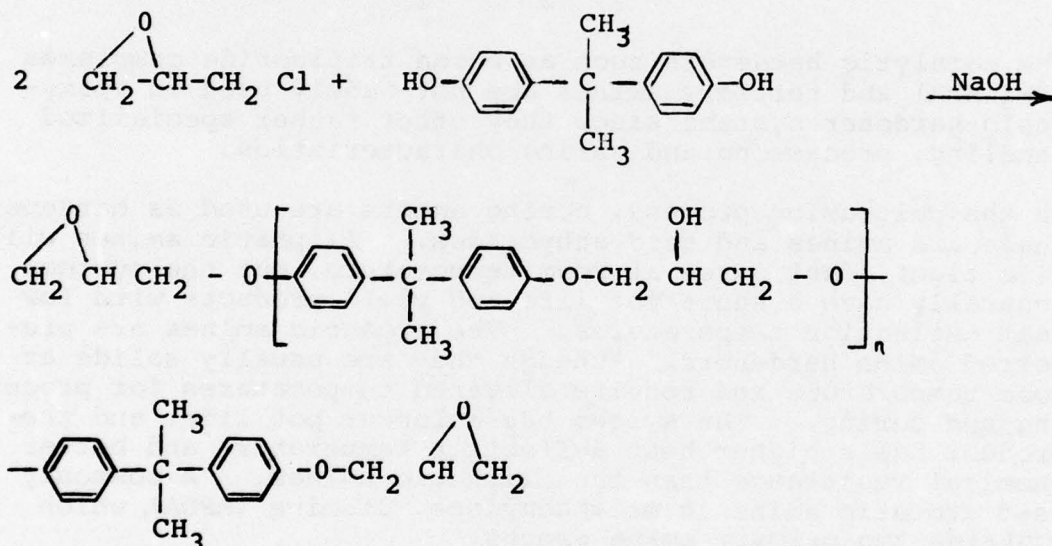
In the above, n determines the molecular weight and the viscosity of the resin. The lower molecular weight products are liquids with an n value of zero. This resin is used in the pultrusion process, since room temperature handling is generally desired. When n is equal to or greater than one, the resins are thermoplastic solids.

An epoxide resin must be converted from a liquid (thermoplastic) state to a tough, hard, solid (thermoset) state by the

101

addition of hardeners, which are chemically reactive toward the epoxide group of the resin.

The hardener may either be a catalyst or a curing agent. A catalyst initiates self-polymerization of the resin, forming direct linkages between the epoxide groups. A curing agent acts as a cross-linking compound, becoming an integral part of the three-dimensional polymer network. The Lewis acid boron trifluoride acts as a catalyst hardener.

$$BF_3 \quad + \quad R-O-CH_2-\overset{\displaystyle O}{\overset{\diagup \diagdown}{CH-CH_2}} \longrightarrow$$

$$\left[R\ O\ CH_2\ \overset{\overset{\ominus}{O}\ BF_3}{\underset{\underset{O}{|}}{CH}}\ CH_2 \right]_n$$

$$R\ O\ CH_2\ CH\ CH_2\overset{\oplus}{\ }\ +$$

The catalytic hardeners such as boron trifluoride complexes (BF$_3$ MEA) and tertiary amines are not widely used in epoxy-resin-hardener systems since they offer rather specialized handling, processing and curing characteristics.
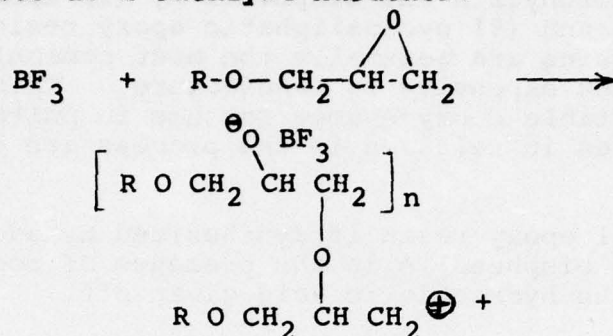
In the pultrusion process, curing agents are used as hardeners; these are amines and acid anhydrides. Aliphatic amines will give tight, fast cures at room temperature, but the systems generally have a short pot life and yield products with low heat deflection temperatures. The aromatic amines are preferred amine hardeners, although they are usually solids at room temperature and require elevated temperatures for processing and curing. The system has a longer pot life, and the product has a higher heat deflection temperature and better chemical resistance than the aliphatic amines. A commonly used aromatic amine is metaphenylene diamine (MPDA), which contains two primary amine groups.

Anhydrides are also used as curing agents. Like aromatic amines, anhydrides require elevated temperature curing; they also yield products with high heat deflection temperatures and good chemical resistance. They provide long pot lives and low exotherms, and anhydrides are available which are low viscosity liquids at room temperature, thus providing low viscosity systems with excellent wetting and processing properties. However, the aromatic amines are preferred since they cause less problems with sticking to die surfaces than the anhydrides.

102

The reaction, when using a primary amine (e.g., MPDA), involves the opening of the epoxide ring through an electronegative attack by nitrogen to form an alcohol without formation of by-products. The hydroxyl group is then available for further reaction, and the secondary amine will react with another epoxy group.

$$\text{I} \quad R - NH_2 \;+\; CH_2 \overset{\displaystyle O}{\overset{\diagup\ \diagdown}{-}} CH - CH_2\, O\, R' \quad \longrightarrow$$

$$R - NH - CH_2\ \underset{\underset{\textstyle OH}{|}}{CH} - O\, R'$$

$$\text{II} \quad \underset{\underset{\textstyle H}{|}}{R-N} - CH_2\ \underset{\underset{\textstyle OH}{|}}{CH} - C \;+\; CH_2 \overset{\displaystyle O}{\overset{\diagup\ \diagdown}{-}} CH - CH_2 - OR \quad \longrightarrow$$

$$R - \underset{\diagdown}{N} - CH_2\ \underset{\underset{\textstyle OH}{|}}{C} - C \sim$$
$$CH_2$$
$$HC-OH$$
$$\diagdown$$
$$C \sim$$
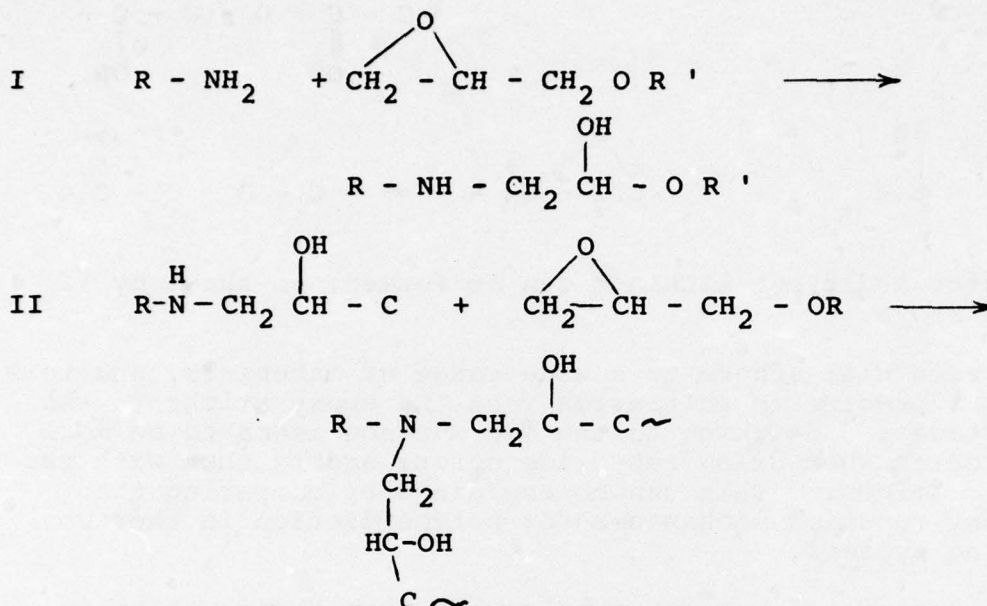
During the reaction, no rearrangement occurs in the backbone of the resin molecule, the reactive epoxide groups being held on the uncured molecule as appendages.

The anhydride (e.g., nodic ethyl anhydride) reaction is initiated by the formation of an ester through the anhydride and a hydroxyl group present on the resin. Water, alcohol, or a tertiary amine can also be used for initiation. Then, the remaining acid group will react with an epoxy group. In addition, self-polymerization of the epoxy group through the epoxy hydroxyl reaction:

$$\text{I} \quad \begin{array}{c} -C - \overset{\displaystyle O}{\overset{\|}{C}} \diagdown \\ | \qquad\quad O \\ -C - C \underset{\displaystyle O}{\diagdown\|} \end{array} \;+\; \sim C - \underset{\underset{\textstyle |}{|}}{\overset{\overset{\textstyle OH}{|}}{C}} - C \sim \quad \longrightarrow \quad \begin{array}{c} -C - \overset{\displaystyle O}{\overset{\|}{C}} - O - C \\ | \\ -C - C - OH \\ \quad\ \ \underset{\displaystyle O}{\|} \end{array}$$

103

II

$$- C - C - O - C \quad + \quad CH_2 - CH - \longrightarrow$$

with the first carbonyl shown as O double bonded, and below:

$$- C - C - OH$$

products:

$$- C - C - O - C$$
$$- C - C - O - C - C -$$

III

$$- C - \quad (OH) \quad + \quad CH_2 - CH - \longrightarrow - C - O - C - C \quad (OH)$$
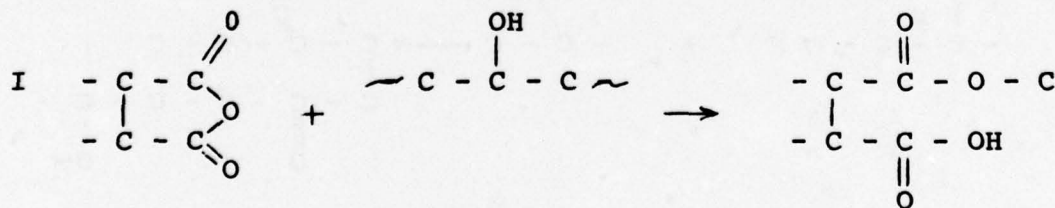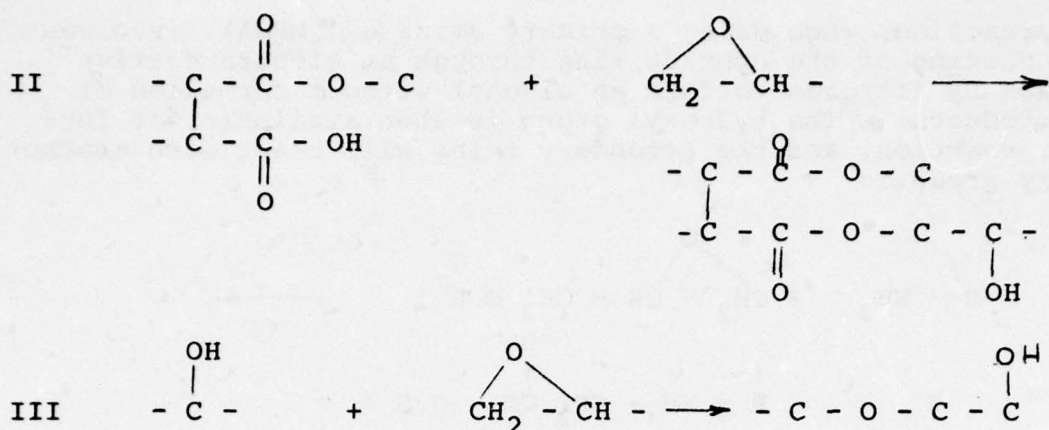
Both ester and ether linkages can be formed, as shown by II and III above.

Epoxy resin will adhere to a wide range of materials, and this becomes a problem in pultrusion when the epoxy sticks to the die surfaces.   Sticking to the die surface seems to be more of a problem when using anhydride curing agents than with the aromatic amines.   This can be explained by comparing the different reaction mechanisms for polymerization in the two hardening systems.

In the presence of oxygen, metals will form oxides at their surface.   The metal oxides will electronegatively attack epoxide groups, creating adhesion at the die surface.

I $\quad M - O \quad + \quad CH_2 - CH - CH_2 - \longrightarrow M - O - CH_2 - CH - \quad (OH)$
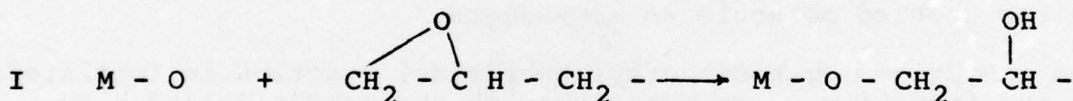
This is similar to the mode of reaction in an acid anhydride, in which acidic hydroxyl groups and secondary alcohols open the epoxy ring through electronegative attack by oxygen:

II

$$- C - C \overset{O}{\Vert}$$
$$- C - C - O H \quad + \quad - C - C \longrightarrow C - C - O - C$$
$$C - C - O - C - C$$

$$\text{III} \quad \underset{\displaystyle -\overset{\displaystyle \overset{OH}{|}}{C}-}{} \quad + \quad \underset{\displaystyle CH_2-CH-}{\overset{\displaystyle O}{\overset{\diagup \diagdown}{}}} \quad \longrightarrow \quad C-O-C-\overset{OH}{\underset{|}{C}}$$
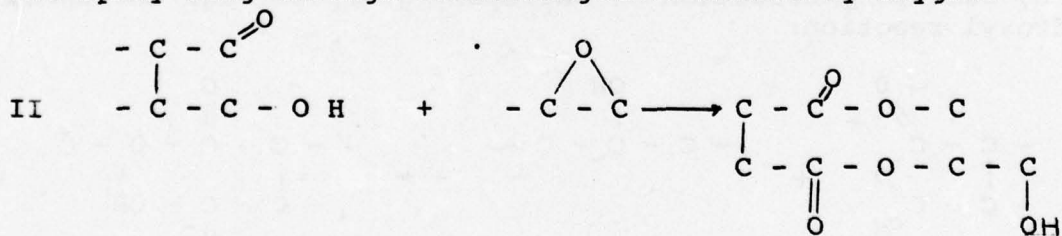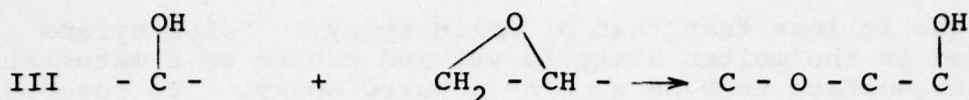
Thus, the metal oxide or metal hydroxyl (M -O-H) group may attack an epoxide ring in a manner similar to the secondary alcohol in the anhydride case.

The same metal oxides are present in the amine hardened system, but the mode of polymerization is opening the epoxide ring. This involves electronegative attack by a nitrogen upon the epoxy ring. Electronegative attack is favored with nitrogen since the lone pair of electrons is more electronegative, and thus more reactive, than the two lone pairs on oxygen. This leads to a greater free energy drop when an epoxide reacts with a nitrogen than with a hydroxyl group, making attack by nitrogen the favored reaction. No chemical reaction is absolute when more than one pathway is possible, but the reaction with the amine group will predominate. Since the anhydride proceeds with electronegative attack by oxygen, side reactions with metal oxides will be more apparent here.

Epoxy resins have adhesion superior to that for polyesters[16]. They will bond to a wide range of materials such as metal, glass, concrete, and plastics. This is due to their wetting and penetrating ability, low viscosity, and the presence of both polar (hydroxyl) and non polar (ether) groups within the molecule, and low degree of volume shrinkage during polymerization (1-6% for epoxy compared with 7-10% for polyester)[17]. The presence of both polar and non polar groups provides outstanding adhesion between dissimilar materials, such as metal and plastic. Wetting allows the epoxy to come into close contact with the surface, and van der Waals[18] law of interaction between molecules comes into action.

Surface energetics in adhesion states that for a liquid to wet and spread on a solid surface, the critical surface tension γ_S of the solids must be greater than the surface tension of the liquid, γ_L, thus

$$\gamma_S > \gamma_L.$$

Metals and metal oxides have surface free energies ranging from 100 to 3,000 ergs/cm^2, and organic liquids, such as epoxy, and water have free energies of less than 100. Whenever oxygen becomes a part of the chemical structure of the surface, polarity and thus surface energy is increased, making wetting by an adhesive such as epoxy easier. The role surface oxidation plays in adhesion may be illustrated by the bonding of liquid epoxy to solid polyethylene. The surface tension of molten

105

polyethylene is less than that of solid epoxy. Polyethylene
may be used in the molten state to wet and adhere to a material
with higher surface tension such as a cured epoxy. To reverse
the situation, i.e., bond liquid epoxy to solid polyethylene,
the surface energy of polyethylene must be made greater than
the surface tension of liquid epoxy. This can be accomplished
by oxidation of the polyethylene surface. This same surface
oxidation occurs in the metal dies used in the pultrusion pro-
cess. Inherent roughness of a surface is also contributory
to adhesion since more surface area is present for molecular
attraction due to van der Waal's forces. Metal dies for pul-
trusion should have highly polished surfaces, and either be
fabricated from or plated with a metal which undergoes minimal
surface oxidation in the presence of air.

APPENDIX B

POLYESTER RESINS - THEIR CHEMISTRY
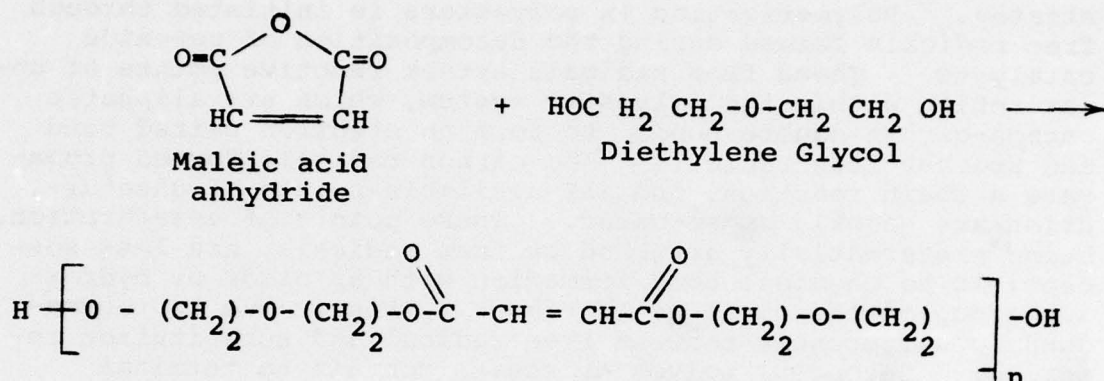## AND APPLICATION TO THE PULTRUSION PROCESS

In the pultrusion process, polyester resin systems give fewer
problems with sticking to metal die surfaces than the epoxy
resins.   This may be attributed to radical chain polymer-
ization in epoxies and the absence in polyesters of reactive
terminal groups which can form chemical bonds to metal sub-
strates.   Polymerization in polyesters is initiated through
free radicals formed during the decomposition of peroxide
catalysts.   These free radicals attack reactive points of un-
saturation within the polyester system, which are aliphatic
carbon-carbon double bonds, to form an electron paired bond
and another free radical.   The carbon radicals formed propa-
gate a chain reaction, and all available points of unsatur-
ation are quickly cross-linked.   These points of unsaturation,
being preferentially attacked by free radicals, are less sus-
ceptible to chemical bond formation with an oxide or hydrox-
ide group on the metal die surface.   This is due to thermo-
dynamic differences between free radical and substitution re-
acting.   Secondly, polyester resins contain no terminal
groups reactive toward chemical groups on metal surfaces.

Chemical Characteristics of Polyesters

The properties of polyester resins may be varied over a wide
range by adjusting the starting materials used for formulation
of the resin.   A vinyl - unsaturated monomer, such as styrene,
is added to the polyester.   This becomes a cross link between
the resin molecules during polymerization and also serves to
lower the viscosity of the system.   The points of unsatur-
ation in the polyester molecules and monomer act as the points
of reactivity by combining together.   Polymerization is ini-
tiated by free radicals formed in peroxide decomposition which
attack reactive carbon-carbon double bonds with formation of
another free radical.   The free radicals formed on polyester
molecules and monomer molecules will attack all other available
points of unsaturation in a chain reaction (propagation).
Free radicals will then combine to end the polymerization re-
action (termination).   Initiation of polymerization propaga-
tion can be accomplished at high temperatures with catalyst
alone, or at room temperature with catalyst and promoter.
The elevated temperature system, which gives a longer pot life,
is better suited to the pultrusion process.

Polyesters used as a matrix for reinforcement with glass fiber
are condensation products of unsaturated dibasic acids, such
as maleic acid anhydride, and dihydric or polyhydric alcohols,

107

such as diethylene glycol. A combination of maleic an-
hydride and diethylene glycol will give a rigid resin when
cured with many points of cross-linking. Resilient and flex-
ible resin systems are obtained by adding saturated aromatic
or aliphatic dibasic acids and by changing the alcohol used.
Fewer points of reactive unsaturation lead to fewer cross-links
between resin and monomer, and this imparts greater flexibility
to the resin. The polyester resin is formed by a condensation
reaction between the dibasic acid and dihydric alcohol as shown
below:

Maleic acid
anhydride

$+$ HOC H_2 CH_2-O-CH_2 CH_2 OH $\longrightarrow$

Diethylene Glycol

In the pultrusion process, a resilient polyester resin is pre-
ferred. This can be manufactured by adding phthalic anhyd-
ride to the above reaction mixture, in a ratio of 1:1 with
maleic anhydride.

Phthalic Anhydride
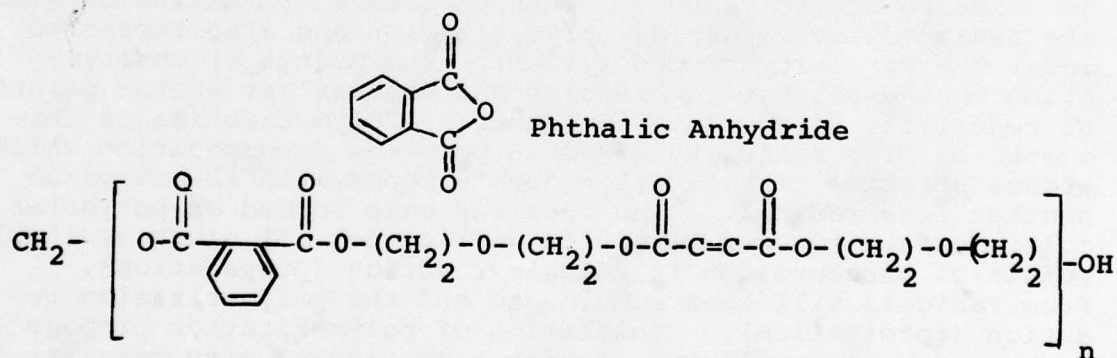
Average polyester molecular weight is determined by the length
of time the glycol and the acid anhydrides are allowed to react
together, and this may be increased by reacting for a longer
time at lower temperatures and by addition of an inhibitor such
as hydroquinone during esterification. A general increase in
molecular weight has a beneficial effect on most of the im-

108

portant properties of the cured resin, such as higher tensile, flexural, and impact strength. The polyester monomer will also attain strength faster during the curing process. When the required molecular weight and viscosity are attained, cooling the reaction mixture will stop condensation. Generally, it is desirable to use as high a molecular weight resin as can be handled without overuse of monomer thinner. Excessive addition of monomer will give a large exothermic heat release, and a composite which is hard, brittle, subject to crazing, and possessing low mechanical and weathering characteristics.

Styrene was originally used as a monomer, and due to economics it has become the most generally used monomeric material. Actually, almost any of the common vinyl unsaturated compounds, which are compatible with the resin and will either homopolymerize or copolymerize with other reactive materials, may be used as a monomer. Between one and two moles of styrene are required for each mole of unsaturated acid in the original ester. Diallyl phthalate may also be added to slow down the cure rate. If a polyester is made from an unsaturated dibasic acid and propylene or diethylene glycol, a highly reactive resin results which forms a rigid thermoset upon curing due to a highly cross-linked structure. This resin is shown above. The addition of a saturated aromatic dibasic acid, such as phthalic anhydride or isophthalic acid, will give the cured polyester more resiliency than the resin with the unsaturated dibasic acid alone. The unsaturation contained within the aromatic ring is not subject to free radical attack, due to the inert nature of the benzene ring, so it does not enter into polymerization. Since there are no points of unsaturation outside the ring for cross-linking, more flexibility of the cured resin is allowed. However, the degree of flexibility allowed by aliphatic chain is not present due to the rigid aromatic ring structure. A flexible resin is manufactured by adding saturated dibasic acids, such as adipic and pimelic acids. Flexibility is imparted by the flexible aliphatic structure between the carboxylic acid groups. The increase in saturation reduces cross-linking, introducing longer chain lengths between cross-linking points. On a molecular level, more bending of the cured resin network is allowed, leading to a more flexible thermoset. Flexibilizers are frequently used in the pultrusion process to minimize thermally induced cracks.

Catalysts for Pultrusion

For pultrusion, a catalyst is desired which will dissociate slowly at room temperature and initiate polymerization with energy supplied through conductive heating or radio frequency radiation. Benzoyl peroxide used at 3% gives a pot

109

life of up to 7 days, and a fast cure within the die when
conductive heating or radio frequency excitation is applied.
Benzoyl peroxide dissociates to form aromatic radicals.
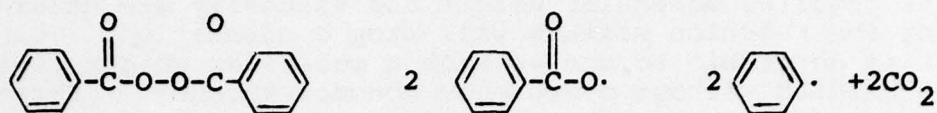
Resistance of Polyester to Environmental Effects

Hydrolysis is the main cause of chemical attack of polyester
resins in an aqueous environment, resulting in chemical decom-
position and resultant rupture of ester linkages in the poly-
mer chain.   Polyester resins possessing increased chemical
resistance and thermal stability (and which also permit
blending with a higher proportion of styrene) may be produced
by condensing isophthalic-based formulations to higher mole-
cular weight.   Such polyester resin systems are now being
used in the pultrusion process.   Another chemically resistant
polyester consists of hydrogenated bisphenol A combined with
maleic, fumaric, and phthalic acids, and propylene glycol.
The increased ring structure helps add chemical resistance; pos-
sibly, absence of tack during curing is imparted by the addi-
tion of bisphenol A.   This resin also exhibits a high heat
deflection temperature, low density, higher viscosity, low
shrinkage during cure, and good adhesion to glass reinforce-
ment, all of which contribute to resistance to hydrolysis.

APPENDIX C

DERIVATION OF STRUCTURAL EFFICIENCY EQUATIONS

Webcore Compression Panel

The mathematical models for the webcore panel considered both overall stability and strength under a uniaxially applied compressive load in the narrow direction of the panel. Overall stability was described using orthotropic sandwich plate theory[15] for a panel with simply supported edges. A maximum strain criterion was used to describe the strength limitations of the panel. The assumptions made were as follows:

1. Plane sections remain plane

2   Only small deflections admissable

3. The panel faces and webs are homogeneous and orthotropic.

4. The thickness of the web and faces are equal

Refer to Figure 43 for a diagram of dimensions used.

The equation for overall stability of an orthotropic sandwich panel as given in Reference 8 page 2.2.2-34 is:

$$N_x = \frac{K\pi^2}{b^2}\sqrt{D_{11}D_{22}} \qquad (C-1)$$

To express (C-1) in the form of a structural efficiency parameter it is convenient to divide both sides of the equation by the panel width b and replace the elastic constants D_{11} and D_{22} with equivalent expressions involving the elastic properties of the material and the panel dimensions. The result is:

$$\frac{N_x}{b} = \frac{a^2}{b\,m^2\pi^2}\left(a_{31} + a'_{31}\right) \qquad (C-2)$$

where a_{31} and a'_{31} are defined on page 2.2.2-34. The expressions for a_{31} and a'_{31} require some modification, however, since the dimensions of the panel are to be given as $\frac{a}{b}$, $\frac{t}{d}$ and $\frac{d}{a}$.

The factor a'_{31} is given in terms of eight expressions. The first simplification, as indicated on page 2.2.2-36, is to set ω equal to zero. Then by noting that $a_{22} \ll a_{13}$, $a_{32} \ll a_{11}$ and $a_{33} \ll a_{21}$, these eight expressions can be reduced to five. Factoring out common constants and substituting into (C-2), the final result is:

111

$$\frac{N_x}{b} = \frac{\pi^2 E_x}{2\,m^2}\left(\frac{1 + {}^t/d}{1 - \nu_{xy}\nu_{yx}}\right)\ \frac{t}{d}\left(\frac{d}{a}\right)^3\left(\frac{a}{b}\right)^5 (A + B)$$

<div align="right">(C-3)</div>

where

$$A = m^4\left(\frac{b}{a}\right)^4 + 2\,m^2\left(\frac{b}{a}\right)^2 n^2\left[\nu_{yx} + \frac{2\,G_{xy}}{E_x}(1 - \nu_{xy}\nu_{yx})\right] + n^4\,\frac{E_y}{E_x}$$

<div align="right">(C-4)</div>

$$B = \frac{\pi^2 E_x}{2}\left(\frac{1 + {}^t/d}{1 - \nu_{xy}\nu_{yx}}\right)\frac{t}{d}\left(\frac{d}{a}\right)^2\left(\frac{a_{21}^2 a_{12} + 2a_{11}a_{21}a_{13} - a_{11}^2 a_{23}}{a_{12}\,a_{23} + a_{13}^2}\right)$$

<div align="right">(C-5)</div>

and

$$a_{11} = -m^3\left(\frac{b}{a}\right) - mn^2\left[\nu_{yx} + \frac{2\,G_{xy}}{E_x}(1 - \nu_{xy}\nu_{yx})\right]$$

<div align="right">(C-6)</div>

$$a_{12} = G'_{cx} + \frac{\pi^2}{2}\left(1 + {}^t/d\right)\frac{t}{d}\left(\frac{d}{a}\right)^2\left[\frac{m^2 E_x}{1 - \nu_{xy}\nu_{yx}} + n^2\left(\frac{a}{b}\right)^2 G_{xy}\right]$$

<div align="right">(C-7)</div>

$$a_{13} = \frac{mn\,\pi^2}{2}\left(1 + {}^t/d\right)\frac{t}{d}\left(\frac{d}{a}\right)^2\left[\frac{E_x\,\nu_{yx}}{1 - \nu_{xy}\nu_{yx}} + G_{xy}\right]$$

<div align="right">(C-8)</div>

$$a_{21} = n^3\,\frac{E_y}{E_x} + nm^2\left(\frac{b}{a}\right)^2\left[\nu_{yx} + \frac{2\,G_{xy}}{E_x}(1 - \nu_{xy}\nu_{yx})\right]$$

<div align="right">(C-9)</div>

$$a_{23} = -\left(\frac{b}{a}\right)^2 G'_{cy} - \frac{\pi^2}{2}\left(1 + {}^t/d\right)\frac{t}{d}\left(\frac{d}{a}\right)^2\left(\frac{b}{a}\right)^2\left[m^2 G_{xy} + n^2\left(\frac{a}{b}\right)^2\frac{E_y}{1 - \nu_{xy}\nu_{yx}}\right]$$

<div align="right">(C-10)</div>

The moduli of the face sheets are calculated by using laminated plate theory, and then with the assumption of homogeneity of the face material, equivalent moduli are calculated. The shear modulus for the webcore in the xz direction, G'_{cx}, was derived by reducing the web shear modulus G'_{xz} by the ratio of web thickness to panel unit width t/w:

$$G'_{cx} = G'_{xz} \left(\frac{t}{w}\right)$$

(C-11)

To relate panel unit width w to web width d, it is assumed that the face between the webs fails by local instability at the same strain as the webs. Using the equation for the local stability failure criteria from Reference 8 page 2.2.1-1, the critical strain for the face is:

$$\varepsilon_f = \frac{G_{xy}}{E_x} \left(\frac{t}{w}\right)^2$$

(C-12)

and for the web:

$$\varepsilon_r = \frac{G'_{cx}}{E'_x} \left(\frac{t}{d}\right)^2$$

(C-13)

Equating (C-12) and (C-13) gives

$$\frac{t}{w} = \sqrt{\frac{E_x}{E'_x} \frac{G'_{cx}}{G_{xy}}} \left(\frac{t}{d}\right)$$

(C-14)

Substituting (C-14) into (C-11) gives:

$$G'_{cx} = \frac{G'^2_{xz}}{G_{xy}} \frac{E_x}{E'_x} \left(\frac{t}{d}\right)^2$$

(C-15)

For G'_{cy}, the web is modelled as a beam clamped at both ends with equal and opposite forces P perpendicular to the beam at each end. Then, the total displacement of the ends is the same as the apparent shear strain giving:

$$\gamma = \frac{Pd^2}{E'_z \, l t}$$

(C-16)

113

The apparent shear stress is:

$$\tau = \frac{P}{\ell t}$$

(C-17)

The shear modulus is τ / γ and by using (C-14) the shear modulus is:

$$G'_{cy} = \left(\frac{t}{d}\right)^3 \sqrt{\frac{G'_{cx}}{G_{xy}} \frac{E_x}{E'_x}} \; E'_z$$

(C-18)

The above expressions for the various moduli are substituted into (C-3). The integer values of m and n are then sought which minimize Nx/b where m and n denote the number of half wavelengths of the buckled panel in the x and y directions respectively.

The structural efficiency parameter for strength can be derived by writing an equation for the running load N_x in terms of strain:

$$N_x = 2 E_x \epsilon_f w \frac{t}{w} + E'_x \epsilon_r d \frac{t}{w}$$

(C-19)

Typically the minimum compressive strain to failure is for a unidirectional layer loaded parallel to the fibers. By assuming that the panel fails when any one layer fails, the unidirectional ultimate strain $\epsilon_{o^\circ}^{cu}$ can be substituted in (C-19). Dividing by b and using (C-14) gives:

$$\frac{N_x}{b} = \epsilon_{o^\circ}^{cu} \frac{t}{d} \frac{d}{b} \left(2 E_x + E'_x \sqrt{\frac{G'_{cx}}{G_{xy}} \frac{E_x}{E'_x}} \right)$$

(C-20)

The weight parameter used was W/ab^2. The total weight of the panel can be expressed as:

$$W = 2 a b t \rho_f + a d t \frac{b}{w} \rho_r$$

(C-21)

Then dividing by ab² and using (C-14), the weight parameter equation becomes:

$$\frac{W}{ab^2} = \frac{t}{d} \frac{d}{b} \left(2 \rho_f + \rho_r \sqrt{\frac{G'_{cx}}{G_{xy}} \frac{E_x}{E'_x}} \right)$$

(C-22)

114

The local instability structural efficiency parameter for the web is:

$$\sigma_{cr} = G'_{cx}\left(\frac{t}{d}\right)^2 = \frac{P_r}{dt}$$

(C-23)

The load carried by the web then becomes:

$$P_r = G'_{cx}\frac{t^3}{d}$$

(C-24)

The total load carried by the panel is:

$$P_{TOTAL} = 2 P_f + P_r\frac{b}{w}$$

(C-25)

The strains in the face and web are given by:

$$\varepsilon_f = \frac{P_f}{bt\,E_x}$$

(C-26)

$$\varepsilon_r = \frac{P_r}{dt\,E'_x}$$

(C-27)

Because of the Assumption 1, ε_f is equal to ε_r. Therefore P_f in terms of P_r is:

$$P_f = P_r\frac{b}{d}\frac{E_x}{E'_x}$$

(C-28)

Substituting (C-24), (C-28) and (C-14) into (C-25) and dividing by b^2 gives:

$$\frac{N_x}{b} = \left(\frac{t}{d}\right)^3\frac{d}{b}G'_{cx}\left(2\frac{E_x}{E'_x} + \sqrt{\frac{G'_{cx}}{G_{xy}}\frac{E'_x}{E'_x}}\right)$$

(C-29)

115

Single-Face Corrugated Compression Panel

An approximate solution to the stability problem for an isotropic plate with longitudinal stringers was obtained by Timoshenko[19] by energy methods. By application of the same method, Lekhnitski [15] also obtained an approximate solution for the case of a special orthotropic plate ($D_{16} = D_{26} = 0$).

The following assumptions were made:

1. Plane sections remain plane

2. Small deflections

3. The plate and stringers are homogeneous and orthotropic

4. The plate deflects and the stringer deflects and twists

Refer to Figure 44 for a diagram of dimensions used. By equating the increased potential energy of the system to the work done by the external forces, an equation for the running load N_x is obtained as:

$$N_x = \sqrt{D_{11} D_{22}} \; \frac{\int_0^b \phi \, dy + b \sum_{i=1}^{N} \left[\gamma_i \beta^4 f^2(\eta_i) + \chi_i \beta^2 f'^2(\eta_i) \right]}{\beta^2 \left[\int_0^b f^2 \, dy + b \sum_{i=1}^{N} \delta_i \, f(\eta_i) \right]} \qquad \text{(C-30)}$$

where

$$\phi = \sqrt{\frac{D_{22}}{D_{11}}} f''^2 - 2\beta^2 \nu_{yx} \sqrt{\frac{D_{11}}{D_{22}}} f f'' + \beta^4 \sqrt{\frac{D_{11}}{D_{22}}} f^2 + \frac{4 D_{66} \beta^2 f'^2}{\sqrt{D_{11} D_{22}}} \qquad \text{(C-31)}$$

$$\delta_i = \frac{A_s E_x'}{b t E_x} \qquad \text{(C-32)}$$

$$\gamma_i = \frac{E_x' I_s}{b \sqrt{D_{11} D_{22}}} \qquad \text{(C-33)}$$

$$\chi_i = \frac{J_s G_{xy}'}{b \sqrt{D_{11} D_{22}}} \qquad \text{(C-34)}$$

116

$$\beta = \frac{m\pi}{a} \,, \quad m = 1, 2, 3 \ldots \ldots \tag{C-35}$$

$$\eta_i = (i - \tfrac{1}{2})(w + d) \tag{C-36}$$

In deriving (C-30), conditions on the supported (loaded) edges and at the end of the stringers were met by assuming that deflections are given by:

$$\Delta = f(y) \sin \beta x \tag{C-37}$$

$$\Delta_i = f(\eta_i) \sin \beta x \tag{C-38}$$

$$\theta = f'(\eta_i) \sin \beta x \tag{C-39}$$

To solve the equation, an expression for functions in the form of a series with undetermined coefficients was introduced as follows:

$$f = \sum_m A_{mn} f_n(y) \tag{C-40}$$

A nontrivial solution requires that the determinant of the coefficients be zero. The smallest root of these homogeneous equations gives the critical load. For a plate with simply supported sides

$$f = \sum_m A_{mn} \sin \frac{n\pi y}{b} = \sum A_{mn} \sin \alpha y \tag{C-41}$$

The above expressions are now substituted into (C-30). Dividing by b^2 the result is:

$$\frac{N_x}{b} = \frac{R + S}{\frac{1}{2} + \frac{A_s E_x'}{bt \, E_x} \sum \sin^2(\alpha \eta)} \tag{C-42}$$

117

where

$$R = \frac{t^3}{24\,b\,(1-\vartheta_{xy}\vartheta_{yx})} \left[E_y \frac{\alpha^4}{\beta^2} + E_x \left[2\vartheta_{yx}\alpha^2 + \beta^2 \right] + 4G_{xy}(1-\vartheta_{xy}\vartheta_{yx})\alpha^2 \right]$$

(C-43)

$$S = \frac{\beta^2 E_x' I_s}{b^2} \sum_{i=1}^{N} \sin(\alpha n_i) + \frac{G_{xy}' J_s \alpha^2}{b^2} \sum_{i=1}^{N} \cos(\alpha n_i)$$

(C-44)

I_s is the moment of inertia of the stringer. According to Reference 19 page 399, this should be taken with respect to the outer surface of the stringer to account for the stiffening effect of the plate.

J_s is a factor like the polar moment of inertia in the torsion of a non-circular shaft. A table of values for various cross-sections can be found in Reference 20. A_s is the *cross-sectional* area of the stringer.

The panel dimensions chosen for this analysis are shown in Figure 44. This results in the following values:

$$I_s = \frac{2}{3} A t d^3$$

(C-45)

$$J_s = d A^3 t^3$$

(C-46)

$$A_s = 3 A t d$$

(C-47)

These expressions are now substituted into (C-42). Furthermore, panel dimensions are rearranged so that all are expressed as t/d, d/b and b/a. The final result is:

$$\frac{N_x}{b} = \frac{K + L}{\frac{1}{2} + 3A \frac{d}{b} \frac{E_x'}{E_x} \sum_{i=1}^{N} \sin^2\left(\frac{n\pi}{b} n_i\right)}$$

(C-48)

where

$$K = \frac{\pi^2}{12(1-\vartheta_{xy}\vartheta_{yx})} \left(\frac{t}{d}\right)^3 \left(\frac{d}{b}\right)^3 \left[\frac{E_y}{2}\left(\frac{a}{b}\right)^2 \frac{n^4}{m^2} + E_x\left(\vartheta_{yx} n^2 + \frac{1}{2}m^2\left(\frac{b}{a}\right)^2\right) + \right.$$
$$\left. 2G_{xy} n^2 (1-\vartheta_{xy}\vartheta_{yx}) \right]$$

(C-49)

$$L = A\pi^2 \left(\frac{t}{d}\right)^3 \left(\frac{d}{b}\right)^4 n^2 \left[\frac{2}{3}\left(\frac{b}{a}\right)^2 \left(\frac{m}{n}\right)^2 \left(\frac{d}{t}\right)^2 E_x' \sum_{i=1}^{N} \sin^2\left(\frac{n\pi}{b}\eta_i\right) +$$

$$A^2 G_{xy}' \sum_{i=1}^{N} \cos^2\left(\frac{n\pi}{b}\eta_i\right)\right] .$$

$$(C-50)$$

The number of stringers N must be expressed in terms of panel dimension ratios and elastic moduli to be used in these expressions. First of all,

$$N = \frac{t}{d+w}$$

$$(C-51)$$

Unit plate width d + w must be related to unit stringer width d. Employing the same method used in deriving (C-14)

$$\epsilon_s = \frac{G_{xy}'}{E_x'}\left(\frac{At}{d}\right)^2$$

$$(C-52)$$

$$\epsilon_p = \frac{G_{xy}}{E_x}\left(\frac{t}{d+w}\right)^2$$

$$(C-53)$$

Equating ϵ_s and ϵ_p

$$\frac{t}{d+w} = \sqrt{\frac{E_x}{E_x'}\frac{G_{xy}'}{G_{xy}}}\,\frac{At}{d}$$

$$(C-54)$$

Substituting (C-54) into (C-51) and simplifying gives

$$N = A\frac{b}{d}\sqrt{\frac{E_x}{E_x'}\frac{G_{xy}'}{G_{xy}}}$$

$$(C-55)$$

This expression for N is substituted into (C-48) and the expression minimized with respect to integer values of m and n. The lowest value is the critical load.

119

The local stability structural efficiency parameter is derived by considering the total load carried by the panel as the sum of the plate load and the stringer loads as follows:

$$P_{TOTAL} = P_P + N P_s \tag{C-56}$$

However, the stringer is made of three equal elements that carry equal loads P_e. The strain in the plate and stringer elements can be expressed as:

$$\varepsilon_P = \frac{P_P}{bt E_x} \tag{C-57}$$

$$\varepsilon_S = \frac{P_e}{d A t E_x'} \tag{C-58}$$

Because of Assumption 1, the strains in the plate and stringer element are equal. Setting $\varepsilon_p = \varepsilon_s$ and solving for P_p gives:

$$P_P = \frac{b}{d} \frac{P_e E_x}{A E_x'} \tag{C-59}$$

The local stability failure criteria for the stringer element is:

$$P_e = A^3 G_{xy}' \frac{t^2}{d} \tag{C-60}$$

Considering that $3P_e = P_s$, substitute (C-59), (C-60) and (C-55) into (C-56). Divide this by b^2 to obtain the local stability parameter as

$$\frac{N_x}{b} = G_{xy}' \left(\frac{t}{d}\right)^3 \frac{d}{b} \left[A^2 \frac{E_x}{E_x'} + 3 A^4 \sqrt{\frac{G_{xy}'}{G_{xy}} \frac{E_x}{E_x'}} \right] \tag{C-61}$$

The ultimate compressive strength parameter is based on the minimum strain criterion. The minimum strain to failure in compression is for a unidirectional layer loaded parallel to the fibers. The loads carried by the plate and a stringer respectively are:

$$P_p = E_x \epsilon_p bt$$

(C-62)

$$P_s = 3 E'_x \epsilon_s dAt$$

(C-63)

Then the total load carried by the panel is:

$$P_{TOTAL} = E_x \epsilon_p bt + 3N E'_x \epsilon_s dAt$$

(C-64)

Again using Assumption 1, the minimum strain criterion is $\epsilon_p = \epsilon_s = \epsilon_{o°}^{cu}$. Dividing (C-64) by b^2 and substituting in (C-55) gives, for the strength parameter

$$\frac{N_x}{b} = \frac{t}{d} \frac{d}{b} \epsilon_{o°}^{cu} \left[E_x + 3 E'_x A^2 \sqrt{\frac{G'_{xy}}{G_{xy}} \frac{E_x}{E'_x}} \right]$$

(C-65)

The weight parameter is W/ab^2. The total weight of the panel can be expressed as:

$$W = abt\rho_p + 3Nadt A \rho_s$$

(C-66)

Dividing by b^2 and using (C-55) the weight parameter equation becomes:

$$\frac{W}{ab} = \frac{t}{d} \frac{d}{b} \left[\rho_p + 3 A^2 \rho_s \sqrt{\frac{G'_{xy}}{G_{xy}} \frac{E_x}{E'_x}} \right]$$

(C-67)

121

LIST OF SYMBOLS

A	constant. Also used to relate stringer thickness to plate thickness
B	constant
D	term in constitutive equation
E	Youngs modulus
G	shear modulus
I	moment of inertia
K,L	constants
N	no. of stringers
P	load
R,S	constants
T	torque
W	structural weight
X	design allowable
a,b,c	dimensions of panel
d	depth (core)
l	unit length (webcore panel)
m,n	half wave lengths
t	panel thickness
w	web or stringer spacing
γ	shear strain
ϵ	strain
ρ	density
τ	shear stress
ν	Poisson's ratio
θ	rotation

Subscripts

cr	critical
cx	web-core
e	P_c refers to 1/3 of total stringer load
f	face (web-core panel)
o	direction with load axis
p	plate (single-face corrugated panel)
r	rib (in web-core panel)
s	stringer (single-face corrugated panel)
x,y,z	coordinate system referred to arbitrary axes
1,2,3	coordinate system referred to material principal axes

Superscripts

'	*web or stringer*
cu	compressive ultimate

14163-76